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- (56) References cited:

EP-A- 0 023 386 EP-A- 0 483 382 EP-A- 0 188 001

WO-A-95/02848

EP-A- 0 572 973

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Description

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BACKGROUND OF THE INVENTION

5 (1) Field of the Invention

[0001] The present invention relates to an optical recording medium, specifically to an optical recording medium which is capable of recording and reproducing in wavelengths of plural lasers.

(2) Description of the Related Art

[0002] CD-R (CD-Recordable) has been developed and proposed as a write-once type optical recording medium which meets compact disk (hereinafter abbreviated as CD) standards (for example, Nikkei Electronics, No. 465, p. 107, January 23, 1989 issue, and OPTICAL DATA STORAGE DIGEST SERIES vol. 1, p. 45, 1989). This CD-R is laminated with a recording layer, a reflective layer and a protective layer in this order on a transparent resin substrate, and irradiation of a laser beam to the above recording layer at a high power causes physical or chemical change on the recording layer and allows information to be recorded thereon in the form of pits. The pits thus formed are irradiated with a laser beam at a low power to detect a change in a reflectance, whereby the information recorded in the pits can be reproduced. Such an optical recording medium uses a near infrared semiconductor laser having a wavelength falling in a range of 770 to 830 nm for recording and reproducing and meets CD standards such as a red book and an orange book, and therefore is characterized by that it is usable for CD players and CD-ROM players.

[0003] In recent years, the development of a semiconductor laser having a wavelength shorter than 770 nm has been advanced, and red semiconductor lasers having wavelengths of 680 nm and 635 nm have been put into practical use (for example, Nikkei Electronics, No. 592, p. 65, October 11, 1993 issue). Shortening of wavelengths of lasers for recording and reproducing can make beam spots small and therefore makes it possible to prepare an optical recording medium capable of recording and reproducing in a high density. Actually investigated are optical recording media having a large capacity and capable of recording information such as dynamic digital images for a long time by using short wavelengths of semiconductor lasers and a data compression technique, and players therefor (for example, Nikkei Electronics, No. 589, p. 55, August 30, 1993 issue and No. 594, p. 169, November 8, 1993 issue). However, even if such high density optical recording media using a red laser and the players therefor have been developed, interchangeability with conventional systems which have already been prevailed widely can not be ignored from a viewpoint of continuity of softwares. That is, required are optical recording media having such interchangeability that recording/reproducing or reproducing is possible with a red laser and recording/reproducing or reproducing is possible as well with a conventional near infrared semiconductor laser having a wavelength of 780 nm.

[0004] Conventional CD and CD-ROM media have less dependence on wavelength in terms of a reflectance and can readily be reproduced with players for high density-media. Meanwhile, since conventional CD-R media use dyes in a recording layer, the optical characteristics depend greatly on wavelengths, and as a result, the reflectances of the CD-R media change to a large extent depending on the wavelengths. For example, while the reflectances to light beam having a wavelength of about 780 nm come up to 65 % or more, the reflectances to red laser having a wavelength selected from a range of 620 to 690 nm are as small as about 10 % and the modulation degrees are small as well because a dye used in a recording layer has a large absorptivity and a small refractive index. In addition, large deformation is observed on the recording wave forms. A reflectance of about 10 % makes it difficult to detect signals, and even if they could be detected by any means, an error rate and a jitter increase, and therefore reproducing with a reproducing player for high density-media will be difficult. Further, there have been created the problems that since the reproduction light is inferior in stability and deterioration is caused merely by reproducing the same truck several times in succession, such conventional CD-R media can not be fit for practical use. Further, caused is low-to-high recording that a recorded part has a larger reflectance than that of a non-recorded part, and the polarity is reverse to that of a conventional CD (high-to-low recording). Accordingly, such CD-R media are not preferred.

[0005] A function-separating recording layer in which a dye layer having a high reflectance and causing no change of the state by heat energy of a laser beam and an organic substance layer having a light absorbing power are laminated in order is proposed in Japanese Patent Application Laid-Open No. 58-112794 (1983) as an example of an optical recording medium provided with two-layered dye. A medium in which a cyanine dye or merocyanine dye having a high reflectance and an organic light absorbing layer are laminated is proposed in Japanese Patent Application Laid-Open No. 60-239948 (1985). Dye recording layers having different optical constants are proposed in Japanese Patent Application Laid-Open No. 63-153192 (1988). A medium in which two kinds of organic dyes having different transmittance and absorptance to a laser beam having a certain wavelength are laminated is proposed in Japanese Patent Application Laid-Open No. 1-110193 (1989). Further, a recording layer comprising an organic two-layer having different transmittance or melting points to a laser beam having a certain wavelength is proposed in Japanese Patent Application Laid-Open No. 1-2001 (1986).

Open No. 4-330649 (1992). However, these proposals are confined to improvement in a reflectance, optical degradation, high sensitivity recording and an error generating rate to a laser beam having a certain wavelength, and they are not optical recording media which are usable for recording and reproducing with plural kinds of laser beams.

[0006] Further, it is proposed in Japanese Patent Application Laid-Open No. 61-74149 (1986) to increase a recording capacity by providing a difference in a depth direction of pits by means of laminating organic dyes having different absorbing wavelengths. However, this also is not an optical recording medium which is usable for recording and reproducing with plural kinds of laser beams.

[0007] On the other hand, media using metal complexes of azo compounds in recording layers are disclosed in Japanese Patent Application Laid-Open No. 62-30090 (1987), Japanese Patent Application Laid-Open No. 63-9577 (1988), Japanese Patent Application Laid-Open No. 63-9578, Japanese Patent Application Laid-Open No. 3-268994 (1991), Japanese Patent Application Laid-Open No. 4-46186 (1992), International Patent Application No. W091/14740, Japanese Patent Application Laid-Open No. 4-308791 (1992), International Patent Application No. W091/18057, International Patent Application No. W091/18950, Japanese Patent Application Laid-Open No. 4-361088 (1992), Japanese Patent Application Laid-Open No. 5-279580 (1993), and Japanese Patent Application Laid-Open No. 6-65514 (1994). However, the media proposed in these publications can not satisfy the orange book and can not be reproduced or recorded and reproduced with beams of 620 to 690 nm.

SUMMARY OF THE INVENTION

[0008] An object of the present invention is to solve the problems described above, that is, to provide an optical recording medium which can be recorded/reproduced or reproduced with light beam having a wavelength selected from wavelengths of 620 to 690 nm and which can be recorded/reproduced or reproduced based on the CD standards as well with light beam having a wavelength selected from conventional wavelengths of 770 to 830 nm and has good recording characteristics.

[0009] Intensive investigations made by the present inventors in order to solve the problems described above have resulted in completing the present invention. That is, the present invention relates to:

(1) an optical recording medium comprising at least a dye layer, a reflective layer and a protective layer provided on a substrate, wherein an azo compound represented by the following Formula (1) or a metal complex thereof having an absorption maximum at a wavelength falling in a range of 450 to 630 nm and a phthalocyanine compound represented by the following Formula (2) having an absorption maximum at a wavelength falling in a range of 680 to 900 nm are contained in said dye layer:

$$R_7 \xrightarrow{Y-N} N = N \xrightarrow{R_3} R_4 R_1$$

$$R_5 \xrightarrow{R_6} R_6$$

$$R_1 \xrightarrow{R_2} R_2$$

$$R_2 \xrightarrow{R_1} R_2$$

wherein R₁ and R₂ represent independently a hydrogen atom, a substituted or unsubstituted alkyl group, a substituted or unsubstituted anyl group, a substituted or unsubstituted analkyl group, or a substituted or unsubstituted alkenyl group; R₃, R₄, R₅ and R₆ represent independently a hydrogen atom, a halogen atom, a hydroxyl group, a carboxyl group, a sulfonic acid group, a sulfonamide group, an amino group, a substituted or unsubstituted alkyl group, a substituted or unsubstituted alkoxyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted acyl group, a substituted or unsubstituted alkylcarboxyl group, a substituted or unsubstituted aralkyl group, a substituted or unsubstituted alkylcarbonylamino group, a substituted or unsubstituted alkylsulfoamino group, a substituted or unsubstituted alkylamino group, a substituted or unsubstituted alkylsulfonyl group, or a substituted or unsubstituted alkenyl group; R_1 and R_4 , R_2 and R_6 , and R_1 and R_2 may form rings via linkage groups; R₇ represents a hydrogen atom, a halogen atom, a hydroxyl group, a carboxyl group, a sulfonic acid group, a sulfonamide group, an amino group, a substituted or unsubstituted alkyl group, a substituted or unsubstituted alkoxyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted acyl group, a substituted or unsubstituted alkylcarboxyl group, a substituted or unsubstituted aralkyl group, a substituted or unsubstituted alkylcarbonylamino group, a substituted or unsubstituted alkylsulfoamino group, a substituted or unsubstituted alkylamino group, a substituted or unsubstituted alkylsulfonyl group, a substituted or unsubstituted alkenyl group, a cyano group, a nitro group, a mercapto group, a thiocyano group, a chlorosulfonic acid group, a substituted or

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unsubstituted alkylthio group, a substituted or unsubstituted alkylazomethine group, or a substituted or unsubstituted alkylaminosulfonyl group; X represents a sulfur atom, or N-R $_8$ (wherein R $_9$ represents a hydrogen atom, a substituted or unsubstituted aryl group, a substituted or unsubstituted aryl group, a substituted aralkyl group, or a substituted or unsubstituted alkenyl group); Y represents a nitrogen atom or C-R $_9$ (wherein R $_9$ is synonymous with those described in R $_7$); provided that when X is a sulfur atom, Y is a nitrogen atom, and when X is N-R $_9$, Y is C-R $_9$:

$$(A_4)_{m4} (Y_8)_{14} = (A_1)_{m1} (Y_8)_{14} = (A_2)_{m2} (Y_8)_{14} = (A_2)_{m2} (Y_8)_{15} = (A_3)_{m3} = (A_2)_{m3} = (A_3)_{m3} = (A_3)_{m3}$$

wherein Y_1 , Y_2 , Y_3 , Y_4 , Y_5 , Y_6 , Y_7 , and Y_8 represent independently a hydrogen atom, a substituted or unsubstituted hydrocarbon group having 1 to 20 carbon atoms, a substituted or unsubstituted alkoxyl group having 1 to 20 carbon atoms, or a substituted or unsubstituted alkylthio group having 1 to 20 carbon atoms; in the respective combinations of Y_1 and Y_2 , Y_3 and Y_4 , Y_5 and Y_6 and Y_7 and Y_8 , they may be combined to form rings when they are adjacent to each other; A_1 , A_2 , A_3 and A_4 represent independently a halogen atom or a nitro group; I_1 , I_2 , I_3 and I_4 represent an integer of 0 to 3; I_1 , I_2 , I_3 , I_4 , I_5 ,

(2) an optical recording medium as described in (1), wherein the compound contained in the dye layer and having an absorption maximum at a wavelength falling in a range of 450 to 630 nm is a metal complex of an azo compound represented by the following Formula (3):

$$R_{11} \xrightarrow{N-N}_{N=N} \xrightarrow{R_{10}}_{N=N} \xrightarrow{R_4}_{R_5} R_1$$
 $R_5 \qquad R_6 \qquad R_2 \qquad (3)$

wherein B_1 , B_2 , B_4 , B_5 , and B_6 are synonymous with those described in Formula (1); B_{10} represents a hydroxyl group or a carboxyl group; and B_{11} represents a hydrogen atom or a halogen atom;

(3) an optical recording medium as described in (1), wherein the dye layer comprises a single layer structure containing a mixture of the azo compound represented by Formula (1) or the metal complex thereof having an absorption maximum at a wavelength falling in a range of 450 to 630 nm and the phthalocyanine compound represented by Formula (2) having an absorption maximum at a wavelength falling in a range of 680 to 900 nm;

(4) an optical recording medium as described in (1), wherein the dye layer comprises a two-layer structure of an optical interference layer containing the azo compound represented by Formula (1) or the metal complex thereof having an absorption maximum at a wavelength falling in a range of 450 to 630 nm and a recording layer containing the phthalocyanine compound represented by Formula (2) having an absorption maximum at a wavelength falling in a range of 680 to 900 nm;

(5) an optical recording medium as described in (4), wherein an equation of 70≤ni x di≤300 applies to light beam having a wavelength used for recording and reproducing, wherein ni represents a real part of a complex refractive index and di represents a layer thickness in the optical interference layer;

(6) an optical recording medium as described in any of (1) to (5), capable of recording and/or reproducing with a laser beam having a wavelength λ_1 selected from wavelengths falling in a range of 770 to 830 and capable of

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recording and/or reproducing as well with a laser beam having a wavelength λ_2 selected from wavelengths falling in a range of 620 to 690;

- (7) an optical recording medium as described in (6), capable of reproducing with a laser beam having the wavelength λ_2 ;
- (8) an optical recording medium as described in (6) or (7), wherein the laser beam having the wavelength λ_1 has a reflectance of 65 % or more, and the laser beam having the wavelength λ_2 has a reflectance of 15 % or more, which are measured through the substrate;
- (9) an optical recording medium as described in (8), wherein the laser beam having the wavelength λ_2 has a reflectance of 20 % or more which is measured through the substrate; and
- (10) a metal complex of an azo compound represented by Formula (4).

$$\begin{bmatrix} R_{11} & N - N \\ S & N = N \end{bmatrix} = N \begin{bmatrix} R_1 \\ R_2 \end{bmatrix} M e^{2\tau}$$
(4)

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wherein R₁ and R₂ represent independently a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 15 carbon atoms, a substituted or unsubstituted aryl group having 6 to 21 carbon atoms, a substituted or unsubstituted aralkyl group having 7 to 22 carbon atoms, or a substituted or unsubstituted alkenyl group 2 to 16 carbon atoms:

R₁₁ represents a hydrogen atom or a halogen atom; Me represents nickel, cobalt, copper, palladium, iron and zinc.

BRIEF DESCRIPTION OF THE DRAWING

- 30 [0010] Fig. 1 is a cross-sectional structural view of the optical recording medium, wherein;
 - 1. substrate,
 - 2. dye layer,
 - 3. reflective layer,
 - 4. protective layer.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

- [0011] The present invention shall concretely be explained below.
- [0012] The optical recording medium of the present invention has a structure in which a dye layer is formed on a substrate and a reflective layer is provided thereon. Further, in order to protect the reflective layer, a protective layer may be provided on the reflective layer, or two sheets of media may be stuck together. However, when the dye layer comprises a two-layer structure of a recording layer and an optical interference layer, a laminating order of the recording layer and the optical interference layer may be reversed, and other layers may be present between the substrate and the dye layer or the optical interference layer or between the recording layer or the optical interference layer and the reflective layer.
 - [0013] The materials for the substrate may be any one as long as it transmits rays having wavelengths for recording and reproducing. Used are, for example, polymer such as polycarbonate resins, vinyl chloride resins, acrylic resins such as polymethylmethacrylate, polystyrene resins and epoxy resins and inorganic materials such as glass. These materials for the substrate are moded into discoid substrates by injection molding. If necessary, guide grooves and pits for showing recording positions are formed on the surface of the substrate in some cases. Such guide grooves and pits are provided preferably in molding the substrate. They can be provided as well by forming a UV-cured resin layer on the substrate. Usually, when the optical recording medium is used for CD, it is a disc having a thickness of about 1.2 mm and a diameter of about 80 to 120 mm, and it has a hole having a diameter of about 15 mm in the center thereof.
 - [0014] In the present invention, the dye layer is provided on the substrate, and the dye layer of the present invention contains a compound represented by Formula (1) or a metal complex thereof having a maximum absorption wavelength (\lambda max) falling in a range of less than 630 nm. Above all, preferred is the compound having a large refractive index and

a small absorbance at wavelengths of 620 to 830 nm. When the dye layer comprises a two-layer structure of the recording layer and the optical interference layer, an attenuation coefficient at λ_1 in the optical interference layer containing a metal complex of the azo compound as a main component is preferably 0.15 or less, and an attenuation coefficient at λ_2 is preferably 0.2 or less.

[0015] The azo compound or the metal complex thereof used in the present invention is the azo compound represented by Formula (1) described above or the metal complex thereof. In the substituents in Formula (1) described above, a halogen atom includes fluorine, chlorine, bromine and iodine, preferably fluorine, chlorine and bromine.

[0016] The number of carbon atoms indicated in the patent claims for the individual meanings of the various symbols there defined corresponds to the total number of carbon atoms including that of possible substituents.

[0017] Whenever appearing in the definition of each one of the individual symbols directed to formulas (1) to (4) the below-mentioned groups can comprise the following meanings.

[0018] An alkyl group is an unsubstitued or substitued alkyl group having 1 to 15 carbon atoms in total. The unsubstituted alkyl group includes a linear, branched or cyclic alkyl group. A substituent of the substituted alkyl group includes a halogen atom, an alkoxyl group, an alkoxyalkoxy group, an alkoxyl group, a hydroxyl group, an acyloxyl group, an acyloxyl group, an acyloxyl group, an acyloxyl group, an alkyl group,

[0019] The above substituents are also the possible substituents of the other listed groups different from an alkyl group, whenever they are substituted.

[0020] A linear or branched alkyl group is an alkyl group having 1 to 15 carbon atoms and includes, considering processability by coating on polycarbonate, acryl, epoxy or polyolefin substrates, methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, t-butyl, n-pentyl, isopentyl, 2-methylbutyl, 1-methylbutyl, neopentyl, 1,2-dimethylpropyl, 1,1-dimethylpropyl, cyclopentyl, n-hexyl, 4-methylpentyl, 3-methylpentyl, 2-methylpentyl, 1-methylpentyl, 3-dimethylbutyl, 2,3-dimethylbutyl, 1,3-dimethylbutyl, 1,2-dimethylbutyl, 1,1-dimethylbutyl, 2-ethylbutyl, 1-ethylbutyl, 1,2-dimethylbutyl, 1,2-dimethylbutyl, 1,2-dimethylbutyl, 1,2-dimethylbutyl, 1,2-dimethylbutyl, 1,2-dimethylbutyl, 1,2-dimethylbutyl, 1,2-dimethylbutyl, 2,5-f-timethylbutyl, 1,2-dimethylbexyl, 2-methylhexyl, 2-methylhexyl, 2-methylhexyl, 2-methylhexyl, 2-methylhexyl, 2-methylhexyl, 2-methylhexyl, 2-methylhexyl, 3-methylhexyl, 4-methylhexyl, 5-methylhexyl, 2,4-dimethylpentyl, n-nonyl, n-decyl, 4-ethyloctyl, 4-ethyl-4,5-dimethylhexyl, n-undecyl, n-dodecyl, 1,3,5,7-tetraethyloctyl, 4-butyloctyl, 6-6-diethyloctyl, n-tridecyl, 6-methyl-4-butyloctyl, n-tetradecyl, n-pentadecyl, 3,5-dimethylhexyl, 2,6-dimethylheptyl, 2,4-dimethylheptyl, 2,2,5,5-tetramethylhexyl, 1-cyclo-pentyl-2,2-dimethylpropyl and 1-cyclohexyl-2,2-dimethylpropyl.

[0021] An alkoxyalkyl group includes groups having 2 to 15 carbon atoms, such as methoxymethyl, ethoxymethyl, propoxymethyl, butoxymethyl, nethoxyethyl, ethoxyethyl, propoxyethyl, butoxyethyl, n-hexyloxyethyl, 4-methylpentyloxyethyl, 1,3-dimethylbutoxyethyl, 2-ethylhexyloxyethyl, n-octyloxyethyl, 3,5,5-trimethylhexyloxyethyl, 2-methyl-1-isopropylpropoxyethyl, 3-methyl-1-isopropylbutoxyethyl, 2-ethoxy-l-methylethyl, 3-methoxybutyl, 3,3,3-trifluoropropoxyethyl and 3,3,3-trichloropropoxyethyl.

[0022] The examples of an alkoxyalkoxyalkyl group include -methoxyethoxyethyl, ethoxyethoxyethyl, propoxyethoxyethyl, butoxyethoxyethyl, 1,2-dimethylpropoxyethoxyethyl, 3-methyl-1-isobutylbutoxyethoxyethyl, 2-methoxy-1-methylethoxyethyl, 2-butoxy-1-methylethoxyethyl, 2-(2'-ethoxy-1'-methylethoxy)-1-methylethyl, 3,3,3-trifluoropropoxyethoxyethyl and 3,3,3-trichloropropoxyethoxyethyl.

[0023] The examples of an alkoxyalkoxyalkoxyalkyl group include methoxyethoxyethoxyethyl, ethoxyethoxyethoxyethyl, butoxyethoxyethyl, butoxyethoxyethyl, 2,2,2-trifluoroethoxyethoxyethoxyethyl and 2,2,2-trichloroethoxyethoxyethoxyethyl.

[0024] The examples of an alkoxycarbonylalkyl group include methoxycarbonylmethyl, ethoxycarbonylmethyl, butoxycarbonylmethyl, methoxycarbonylethyl, ethoxycarbonylethyl, butoxycarbonylethyl, 2,2,3,3-tetrafluoropropoxycarbonylmethyl and 2,2,3,3-tetrafluoropropoxycarbonylmethyl.

[0025] The examples of an alkoxycarbonyloxyalkyl group include methoxycarbonyloxyethyl, ethoxycarbonyloxyethyl, butoxycarbonyloxyethyl, 2,2,2-trifluoroethoxycarbonyloxyethyl and 2,2,2-trichloroethoxycarbonyloxyethyl.

[0026] The examples of an alkoxyalkoxycarbonyloxyalkyl group includes methoxyethoxycarbonyloxyethyl, ethoxyethoxycarbonyloxyethyl, butoxyethoxycarbonyloxyethyl, 2,2,2-trifluoroethoxyethoxycarbonyloxyethyl and 2,2,2-trichloroethoxyethoxycarbonyloxyethyl.

[0027] The examples of a hydroxyalkyl group include 2-hydroxyethyl, 4-hydroxybutyl, 2-hydroxy-3-methoxypropyl, 2-hydroxy-3-chloropropyl, 2-hydroxy-3-ethoxypropyl, 3-butoxy-2-hydroxypropyl, 2-hydroxy-3-phenoxypropyl, 2-hydroxypropyl and 2-hydroxybutyl.

[0028] The examples of a hydroxyalkoxyalkyl group include hydroxyethoxyethyl, 2-(2'-hydroxy-1'-methylethoxy)-1-methylethyl, 2-(3'-fluoro-2'-hydroxypropoxy)ethyl and 2-(3'-chloro-2'-hydroxypropoxy)ethyl.

[0029] The examples of a hydroxyalkoxyalkoxyalkyl group include hydroxyethoxyethoxyethyl, [2'-(2"-hydroxy-1"-methylethoxy)-1'-methylethoxy]ethoxyethyl, [2'-(2"-fluoro-1"-hydroxyethoxy)-1'-methylethoxy]ethoxyethyl, [2'-(2"-chloro-1"-hydroxyethoxy)-1'-methylethoxy]ethoxyethyl,

- [0030] The examples of a cyanoalkyl group include 2-cyanoethyl, 2-cyanopropyl, 2-cyanobutyl, 4-cyanobutyl, 2-cyano-3-methoxypropyl, 2-cyano-3-chloropropyl, 2-cyano-3-ethoxypropyl, 3-butoxy-2-cyanopropyl and 2-cyano-3-phenoxypropyl.
- [0031] The examples of an acyloxyalkyl group include acetoxyethyl, propionyloxyethyl, butyryloxyethyl, valeryloxyethyl, 1-ethylpentylcarbonyloxyethyl, 2,4,4-trimethylpentylcarbonyloxyethyl, 3-fluoro-butyryloxyethyl and 3-chlorobutyryloxyethyl.
 - [0032] The examples of an acyloxyalkoxyalkyl group include acetoxyethoxyethyl, propionyloxyethoxyethyl, valeryloxyethoxyethyl, 1-ethylpentylcarbonyloxyethoxyethyl, 2.4,4-trimethylpentylcarbonyloxyethoxyethyl, 2-fluoropropionyloxyethoxyethyl, and 2-chloropropionyloxyethoxyethyl.
- 10 [0033] The examples of an acyloxyalkoxyalkoxyalkyl group include acetoxyethoxyethoxyethyl, propionyloxyethyl.
 - [0034] The examples of a halogenated alkyl group include chloromethyl, chloroethyl, 2,2,2-trifluoroethyl, trifluoromethyl, bromomethyl and iodomethyl.
 - [0035] The examples of a sulfonalkyl group include sulfonmethyl, sulfonethyl and sulfonpropyl.
 - [0036] The examples of an alkylcarbonylaminoalkyl group include methylcarbonylaminoethyl, ethylcarbonylaminoethyl, propylcarbonylaminoethyl, cyclohexylcarbonylaminoethyl and succiniminoethyl.
 - [0037] The examples of an alkylsulfoaminoalkyl group include methylsulfoaminoethyl, ethylsulfoaminoethyl and propylsulfoaminoethyl.
- 20 [0038] The examples of a sulfonamidealkyl group include sulfonamidemethyl, sulfonamidethyl and sulfonamidepropyl.
 - [0039] The examples of an alkylaminoalkyl group include N-methylaminomethyl, N,N-dimethylaminomethyl, N,N-dipropylaminomethyl and N,N-dibutylaminomethyl.
 - [0040] The examples of an aminoalkyl group include aminomethyl, aminoethyl and aminopropyl.
- 25 [0041] The examples of an alkylsulfonalkyl group include methylsulfonmethyl, ethylsulfonmethyl, butylsulfonmethyl, methylsulfonethyl, ethylsulfonethyl, butylsulfonethyl, 2,2,3,3-tetrafluoropropylsulfonmethyl and 2,2,3,3-tetrachloropropylsulfonmethyl.
 - [0042] The examples of a substituted or unsubstituted alkoxyl group are alkoxyl groups having the same substituents as the alkyl groups given above have, and include preferably lower alkoxyl groups such as methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, isobutoxy, sec-butoxy, t-butoxy, n-pentyloxy, isopentyloxy, neopentyloxy and 2-methylbutoxy.
 - [0043] The examples of a substituted or unsubstituted aryl group are aryl groups having the same substituted as the alkyl groups given above have, and include preferably phenyl, nitrophenyl, cyanophenyl, hydroxyphenyl, methylphenyl, trifluoromethylphenyl, naphthyl, nitronaphthyl, cyanonaphthyl, hydroxynaphthyl, methylnaphthyl and trifluoromethylnaphthyl.
- [0044] The examples of a substituted or unsubstituted acyl group are acyl groups having the same substituents as the alkyl groups given above have, and include preferably formyl, acetyl, ethylcarbonyl, n-propylcarbonyl, isoputylcarbonyl, sec-butylcarbonyl, t-butylcarbonyl, n-pentylcarbonyl, isopentylcarbonyl, neopentylcarbonyl, 2-methylbutylcarbonyl and nitrobenzylcarbonyl.
- [0045] The examples of a substituted or unsubstituted alkylcarboxyl group are alkylcarboxyl groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylcarboxyl groups such as methylcarboxyl, ethylcarboxyl, n-propylcarboxyl, isopropylcarboxyl, n-butylcarboxyl, isobutylcarboxyl, sec-butylcarboxyl, t-butylcarboxyl, n-pentylcarboxyl, isopentylcarboxyl, neopentylcarboxyl and 2-methylbutylcarboxyl.
 - [0046] The examples of a substituted or unsubstituted aralkyl group are aralkyl groups having the same substituents as the alkyl groups given above have, and include preferably benzyl, nitrobenzyl, cyanobenzyl, hydroxybenzyl, methylbenzyl, trifluoromethylbenzyl, naphthylmethyl, nitronaphthylmethyl, cyanonaphthylmethyl, hydroxynaphthylmethyl, methylnaphthylmethyl and trifluoromethylnaphthylmethyl.
 - [0047] The examples of a substituted or unsubstituted alkylcarbonylamino group are alkylcarbonylamino groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylcarbonylamino groups such as acetylamino, ethylcarbonylamino, n-propylcarbonylamino, isopropylcarbonylamino, n-butylcarbonylamino, isopropylcarbonylamino, i
 - [0048] The examples of a substituted or unsubstituted alkylsulfoamino group are alkylsulfoamino groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylsulfoamino groups such as methylsulfoamino, ethylsulfoamino, n-propylsulfoamino, isopropylsulfoamino, n-butylsulfoamino, isobutylsulfoamino, sec-butylsulfoamino, t-butylsulfoamino, n-pentylsulfoamino, isopentylsulfoamino, neopentylsulfoamino, 2-methylbutylsulfoamino and cyclohexylsulfoamino.
 - [0049] The examples of a substituted or unsubstituted alkylamino group are alkylamino groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylamino groups such as N-meth-

ylamino, N,N-dimethylamino, N,N-diethylamino, N,N-dipropylamino and N,N-dibutylamino.

[0050] The examples of a substituted or unsubstituted alkylsulfonyl group are alkylsulfonyl groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylsulfonyl groups such as methylsulfonyl, ethylsulfonyl, n-propylsulfonyl, isopropylsulfonyl, n-butylsulfonyl, isobutylsulfonyl, sec-butylsulfonyl, t-butylsulfonyl, n-pentylsulfonyl, isopentylsulfonyl, neopentylsulfonyl, 2-methylbutylsulfonyl, 2-hydroxyethylsulfonyl and 2-cyanoethylsulfonyl.

[0051] The examples of a substituted or unsubstituted alkenyl group are alkenyl groups having the same substituents as the alkenyl groups given above have, and include preferably lower alkenyl groups such as propenyl, 1-butenyl, isobutenyl, 1-pentenyl, 2-methyl-1-butenyl, 2-methyl-1-butenyl, 2-methyl-2-butenyl, 2,2-dicyanovinyl, 2-cyano-2-methylcarboxylvinyl and 2-cyano-2-methylsulfonvinyl.

[0052] The examples of a substituted or unsubstituted alkylthio group are alkylthio groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylthio groups such as methylthio, ethylthio, n-propylthio, isopropylthio, n-butylthio, isobutylthio, sec-butylthio, t-butylthio, n-pentylshio, isopentylthio, neopentylthio, 2-methylbutylthio and methylcarboxylethylthio.

[0053] The examples of a substituted or unsubstituted alkylazomethine group are alkylazomethine groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylazomethine groups such as methylazomethine, ethylazomethine, n-propylazomethine, isopropylazomethine, n-butylazomethine, isoputylazomethine, sec-butylazomethine, t-butylazomethine, n-pentylazomethine, isopentylazomethine, n-pentylazomethine, 2-methylbutylazomethine and hydroxyethylazomethine.

[0054] The examples of a substituted or unsubstituted alkylaminosulfonyl group are alkylaminosulfonyl groups having the same substituents as the alkyl groups given above have, and include preferably lower alkylaminosulfonyl groups such as N-methylaminosulfonyl, N-ethylaminosulfonyl, N-(n-propyl)aminosulfonyl, N-(isopropyl)aminosulfonyl, N-(n-pentyl)aminosulfonyl, N-(isopotyl)aminosulfonyl, N-(isopotyl)aminosulfonyl, N-(n-pentyl)aminosulfonyl, N-(1-pentyl)aminosulfonyl, N-(2-hydroxyethyl)aminosulfonyl and N-(2-cyanoethyl)aminosulfonyl.

[0055] The examples of the rings formed by combining R₁ with R₄ and R₂ with R₆ via linkage groups include -CH₂CH₂-, -CH₂CH₂CH₂-, -CH₂CH₂-, -CH₂-, -CH₂CH₂-, -CH₂CH₂-, -CH₂-, -CH₂CH₂-,

[0056] The examples of the rings formed by combining R₁ with R₂ via a linkage group include -CH₂CH₂OCH₂CH₂-, -CH₂CH₂NHCH₂CH₂-, -CH₂C(=O)OC(=O)CH₂-, -CH₂C(=O)NHC(=O)CH₂-, -CH₂C(=O)N (CH₃)C(=O)CH₂- and -CH₂CH₂CH₂CH₂CH₂-.

[0057] The compound of the present invention represented by Formula (1) can be produced in the following manner by known methods. That is, an amine component represented by the following Formula (4) is turned into an azo component, which is added to a solution of a coupling component represented by the following Formula (5) to carry out a coupling reaction, whereby the azo compound represented by Formula (1) is obtained:

$$H_7 = \begin{array}{c} Y - N \\ N \\ N \\ N \end{array}$$
 (4)

$$\begin{array}{c}
R_3 \\
R_5
\end{array}$$

$$\begin{array}{c}
R_4 \\
R_2
\end{array}$$

$$\begin{array}{c}
R_1 \\
R_2
\end{array}$$

$$\begin{array}{c}
(5)
\end{array}$$

wherein R₁ to R₇, X and Y are synonymous with those described in Formula (1).

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[0058] In the present invention, the metal complex of the azo compound described above can be produced by known methods, for example, a method described in Furukawa, Analytica Chimica Acta 140 (1982), 281 to 289. Preferred as a metal for forming the metal complex of the azo compound are, for example, metals such as nickel, cobalt, iron, ruthenium, rhodium, palladium, copper, osmium, iridium, platinum, zinc and magnesium, and more preferred are nickel, cobalt, copper, palladium, iron and zinc. They are used in the forms of acetates, halides and BF₄ salts to form complexes in which the metals are coordinated to the azo compounds in the forms of Ni²+, Co²+, Co³+, Cu²+, Pd²+, Fe²+, Fe³+

and Zn²⁺. The metal complexes of the azo compounds may be used alone or in a mixture of the plural compounds. [0059] The examples of the azo compound of the present invention represented by Formula (1) and the metal complex thereof are shown in Table 1.



55	45 50	35 40		30	25	20		10		5
T.	Table 1									
Compound	R ₁	P ₂	æ	ď	ዊ	æ	4	×	>	Metal
1-1	сы гон	CH2CH2CH	Ŧ	. L	I	 x	ર્દે	N-CH2CH3	3	
1.2	CH2CH2COCH3	. chicacini	I	I	I	I	ર્ક	₩ -₩-₩	So So	•
1-3	G, G, G, G,	CH2CH2CH3	=	I	x	I	*	N-CH2CH3	CSCH3	•
1-4	CH3COCH2CF2CHF2	ರ ₁₃ ಯರರ್ಗ್ನರ್್	x	I	I	×	ફ	N-CH	ફુ	
1-5	G42G43	6 36	I	r	¥	I	ર્દ	N-CH3	3	•
1-6	д ,	Grand Grand	I	I	₹	x	ક	K C F	ਲ	
1-7	දි දී	CH2CH2N4CCH3	r	r	ቼ	I	ર્સ	¥	3	,
1.8	G12012013	CH2CH2CCCCHCH3	I	I	₹	I	x	Q	CCCCH	•
1-9	CH2CH2CH3	ಈ <u>,</u> ಈ,ಈ,ಈ,	I	œ42¢43	₹	I	Ξ_	N-CH#CHCH	N-CHachong Coochs	•
1-10	ਨੂੰ ਨੂੰ	G42G43 ·	I	I	N-HOOCH ₃	x	<u>.</u>	N-CH2CH3	ş	•
1-11	CH2CH2CH3	CH2CH2CH3	x	x	NHCOCH ₃	I	ર્ફ	Ŧ	c∞∞ದಗ್ಕಳು	٠,
1-12	्र ह	\rightarrow \(\vec{\pi}{\pi} \)	x	I	NHCOCH ₃	I	ર્દ	N-CH3	CCONH ₂	•
1-13	CH2CH2CH(CH3)	GH2CH2CH(CH3)2 CH2CH2CCH2)2	T z	I	NHCOCH3	I	I	N-CH ₃	COOH	•
1-14	ಆ್ತ್ರಚ್ಚುಚ್ಯ	<u> </u>	I	x	NHCOCH3	I	Ą	Q) 0-0-0	
1-15	Q ₂ Q ₃	Q242A	x	8	NHCOCH	I	ਤੰ	ĭ	₽ ಡಿಯ	·
1-16	Q2,Q2,Q3	ರ್ಷಚ್ಚರ್ಭಚ್ಯ	x	I	NHCO—(H)	×	ર્દ	¥	CNO2	

22	45 50	35 40		30	25	20	15	10	5	
Te	Table 1 (continued)	d)								
Compound	P,	Pè	ዋ	ď	ጜ	8	4	×	>	Metal
1-17	CH1COCCH2CF2CHF2	CH3COCH2CF2CHF2	Ξ	Ξ	N-CO-N	Ť	x	N-CH2CH3	3	•
1-18	G4;G4;G4;	ಆಸ್ತಿ ಆಸ್ತರ್ಣ	I	x	(Poote)	I	ર્દ	Ŷ	S	•
1-19	Q2,Q42,Q4	СМ2СМ2СМ	I	8	NHOO-	x	ર્દ	N-CH ₂ Q43	S CS	٠
1-20	GH2CH2CH3	നൂ മൂ മൂ	I	I	NHOOCH, CH2G H	Σ Ö	x	N. 042	CCCCCH ₃	•
1-21	Combined with R,	Combined with R.	Ξ.	(CH ₂) ₃	۵	(CH ₂) ₃	r	N-CH3	CN02	•
1-22	Combined with R	Combined with R	I	CH2COCH2	ર્દ	⇔್ತ∞⇔, ⇔₂⊶,	, 942943	Ī	SOS	•
1-23	Combined with R,	CH2CH2CH2	I	I	NHCOCH ₃	I	CH2CH3	¥	CCCOCH ₃	,
1-24	, d1,d2,d1,d1,	x	I	I	8 H	I	ō	¥	C.SO ₃ H	•
1-25		I	x	I	80° H€	I	ર્દ	₹	C.SO2NH2	•
1-26	G12G2G13		x	I	802NH2	I	ਰੰ 	Ŧ	C-NH2	•
1-27	CH#CHCH ₃	CH-CHCH3	x	I	NH2	¥	ર્કે	Ŧ	COCH	•
1-28	CH2CH2CH3	ರ್ಚಿದ್ದರ್ಭ	I	I	COOCH ₃	I	ર્દ	¥	Q	•
1-29	⇔²⇔²⇔₃	ಈ2⇔2⇔3 ಈ2	I	r	NHSO ₂ CH ₃	x	ર્જે	¥	Ç ğ	•
1-30	ರ್ಯಭ್ಯಭ್ಯ	ಈ <u>,</u> ಈ,ಈ,	I	I	0	I	ર્સ	Ī	CNHOOCH	•
1-31	ರ್ಚಿದ್ಯಾರ್ಚ್ಕ	<u> </u>	x	I	8	x	x	Ī	C-NHSO ₂ CH ₃	•
1-32	G42G42G43	ಈ, ಭ್ಯಚ್ಚಿ	x	I	Q Z	x	Ι	H	C-N(CH ₃) ₂	

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	· R ₆ R ₇ X Y Metal	2 H H N.CH3 C.CH.CH3 -	H. H N.CH, C.SH	13 H H CSCN	H OH NGH3 CSO2A	H H C-CH-NCH2OH2OH -	
	B	NH(CH ₃) ₂	80 ₂ CH ₃	CH=CHCH3	x	.	;
	G,	x	Ι	I	I	I	:
	æ	I	x	I	I	I	:
1)	P _B	CH2CH2CH3	₩²₩°	G42G4	CH2CH2CH	₩²₩²₩	2
Table 1 (continued)	R,	ch2ch2ch3	CH2CH2CH	A2424	Q2Q2Q4	H2, CH2, CH	
Tabl	Compound	1-33	1-34	1-35	1-36	1-37	1

	1	1																1
5		Metal		•	•	•			•	•	•		•		•	•	•	
		>	z	z	z	z	z	z	z	z	z	z	z	Z	z	z	z	z
10																		
		×	S	Ø	ဟ	တ	ဟ	Ø	တ	ဟ	Ø	Ø	ທ	Ø	တ	ဟ	S	S
15		æ	ર્ફ	ģ	ษ์	I	·≖	I	ä	Ř	8	8	ă	ă	ă	ĕ	r	$\infty_2 cH_3$
20		CEP	I	I	I	.¤	I	I	I	x	I	I	x	I	I	x	I	I
25		5	x	x	I	I	₹	₹	8	ቼ	₹	₹	NHCOCH3	NHCOCH ₃	NHCOCH ₃	NHOOCH ₃	NHCOCH ₃	NHCOCH,
30		ď	S F	x	I	x	I	x	x	I	x	OCH2CH3	I	I	I	x	I	OCH3
		æ	I	x	I.	I	x	x	I	r	x	x	I	r	I	x	x	Ξ
35				ž.	ę.	OF.		ਨੂੰ		£	ş	₽.		ર્સ		-(CH3)	·	,
40	d)	ď.	CH2CH2CH	CH2CH2COCH3	CH2CH2CH3	CH3COCH2CF2CHF2	GH2GH3	ರ್ಚಿಚ್ಚರ್ಚಿಯ	ઌ૾ૢ૽ૡૢૺૺૺૺૺૺૺૺૺૺૺૺૺૺૺૺૺૺ	O CH2CH2NHCOCH3	CH2CH2COCHCH3	9 ,99,99,99,9	ਲੰ ਲੋ	<u> </u>	\$ 50 m	2 ೧೫,೦೦೫,೦೦೫,೦೧	8,8,8,8	94949 94949
45	Table 1 (continued)	Ę	CH2CH2CH	ಡ್ತುದ್ಯಾಯದ್ಯ	ch2ch2ch3	ರ ₁ ಯ೦೦೫,⊄₂೦№2	o _z o ₃	ಈ₂ಈ₂ಈ₃	ch2ch3	ದ್ಯಾ ರ್ತ್ಕ	ಈ,ಈ,ಈ <u>,</u> ಈ,	ಈ ₂ ಈ2ಈ3	onzon,	- CH2 CH2 CH3	O F	CH2CH2CH(CH3)2 CH2CH2CCH(CH3)2	**************************************	CH2CH3
50	Tab1	Compound	1-39	1-40	1-41	1-42 0	1-43	1-44	1-45	1-46	1.47	1-48	1-49	1-50	1.5.1	1-52 a	1-53	

5		γ Metal	z	, z	z	z	, z	. z		, z		, z	. 2	·	·	·		
10		×	S	S	S	S	ဟ	ဟ	တ	Ø	တ	ဟ	ဟ	υ	တ	ဟ	ဟ	S
15		R,	Q ₃	SCH2C42COCH3	I	Ξ.	CH=NCH2CH2OH	ă	x	=	22	à	מ	င်	N(CH(CH ₃) ₂) ₂	Ξ	Ŧ	I
20		g.	æ	r	x	x	I	I	r	(CH ₂) ₃	CH2COCH2	x	r	I	I	x	I	I
25		æ	NHCO—(H)	NHCO-H)	NHCO-H	NHCO-(H)	NHCO	NH00-	NHOOCH, CH2C	ت	ર્ફ	NHCOCH ₃	5	ž	8	NHOO-H	NHCOCI (3	1 000
30		رځي	Ξ	I	I	8 F	r	£	I	(CI43)2	CH2COCH2	x	r	I	Ξ	I	8 8	I
		æ	Ξ	I	x	I	I	Ξ	Ξ	x	I	I	I	I	I	I	I	I
35 40	d)	R	HO2HO2HO	CH2CH2CH3	CHOCOCH2CF2CHF2	Q ₂ Q ₃	ದ್ಯುದ್ಭುದ್ಯ	H02H02H0	⇔. ಆ₂⇔.⇔.	Combined with R	Combined with R	a12042042	GH2CH2CH3	a1,cH2CH2CH3	65°G3	H02H02H0	Q42Q43	<u> </u>
45	Table 1 (continued)	R ₁	CH2CH2CH	G12G12G13	a 1,0000H20F204F2	a 1, a 1,	012012013	a _{1,} a _{1,} o ₁	042042043	Combined with R	Combined with R.	Combined with R,	042042013	ಈ, ಭ್ಯಭ್ಯಭ್ಯ	al ₂ aH ₃	, ज्यु	GH ₂ GH ₃	a12a12a13
50	Ta}	Compound	1-55	1-56	1-57	1-58	1-59	1-60	1-61	1.62	1-63	1-64	1-65	1-66	1-67	1-68	1.69	1-70

Tak	Table 1 (continued)	. (pan				•				
Compound	, B,	ď	₆	12	ዳ	a ^B	P,	×	>	Metal
1.87	CH2CH3	ದ್ಯಿದ್ಯ	Ξ	X	M+CO-H	r	80°C	S	z	
1-88	CH ₂ CH ₃	CH2CH3	x	Ξ	NHCO—(H)	x	SO2NHCH ₃	ဟ	z	

	45 50	40	30 35	25		20		15	10	5
B	Table I (continued)	Jed)								
Compound	ፎ	R	Ъ	ď	망	d ^a	В,	×	>	Metal
1-89	012012012013	ರ್ಬಿದ್ಬಬ್ಬರು	₹	Ι	r	I	ာ် -	N-CH2Q43	S	Z
1-90	a12a120012013	G12C42CC43CH3	NHCOCH ₃	I	I	8	g,	N-O12	CCN	Z
1-91	042	₽	NHCOCH3	⊗ H¢	=	80 ₃ H	=	N-CH ₂ CH ₃	C-SCH ₃	P
1-92	םיו ₂ כא ₂ סכא ₂ כא(כא	01204200H2CH(CH3)20120H2CH(CH3)2	NHCOCH	x	I	X	ğ	N-CH3	O	Ż
1.93	012013	Q42,Q43	ĕ	I	I	I	ર્દ	N-CH ₃	Son	Ż
1-94	a12012013	ರ್ಬಿರ್ಬಿರ್ಬಿ	₹	I	x	I	ક	N-CH ₃	8	Z
1-95	G12CH(CH3)2	Q12CH(CH3)2	NHCOCH ₃	٥	x	Ö	ર્દ	ž	S	Zn
1.96	G12Q12Q12	ರ್.ದ.ದ.ದ್ಯ	₹	r	×	I	=		ссоосн	ž
1.97	G12012013	<u> </u>	₹	x	I	I	¥	N-CH=CHCH3	C.CCOCH ₃	Z
1.98	Q-12Q-12Q-1	91,04,0H	NHCO-(H)	Ξ	I	₹	ĕ	N-CH2QH3	CCN	Pd
1-99	ಅ್ಯಜ್ಯಂಚ್ಯಂಚ	CH2CH2CCH2CH	NHCOCH ₃	I	x	NHCOCH3	ફ	Ŧ.	C-COOCH Is	Z
1-100	.012013	C42C43	₹	Ξ	x	x	ğ	N-CH3	C-CONH ₂	Ī
1-101	C42C12NISO2C43	CH2CH2NHSO2CH3	ĕ	x	±	Ξ	I	N-Cit ₃	CCOOH	Z
1-102	012012012013	ರ್ನಿದ್ಬರ್ಭರು	₹	Ξ	I	I	ည်	N-CON	89	Ž
1-103	a12a12a12a13	CH2CH2C12CH3	ĕ	I	I	I	ğ	¥	(<u>)</u> .2000	Z
1-104	CH ₂ CH ₃	ರ್ಡಿಚ್	COOH	H	Ξ.	I	Q.	NH	CNO2	Z

Table 1 (continued)		
1 (continu	7	ב פ
-	1	3
Table 1	Ċ	֚֭֭֭֭֭֭֭֭֭֭֭֭֭֭֭֭֭֭֭֭֡֝֝
Tab	9	י
	E	3

				1	1					
Compound	E.	£	ع	2	æ	20	£.	×	>	Metal
1-105	0420011201200H3	CH2 CCH2 CCH3	80 HO	Ξ	I	=	I	N-CH ₂ CH ₃	CC	ž
1-106	a f2 a 42 NHCOCH i3	CH2CH2NHCOCH3	800 HOO	80 HO	I	π	o G		CCC	Z
1-107	ρ. Ο (5α	120CH2CH(CH3)2CH2CH2CH(CH3)2	80°	ж. Ж.	I	SO ₃ H	ğ	N-CH ₂ OH ₃	SCN	Z
1-108	G12OH3	. 04 ₂ 04 ₃	NHSO ₂ CH ₃	S.	I	I	=	NCH2	и-сн ₂ —() ссоосн ₂ сн ₃	Z
1-109	012012012013	ರ್ಬಿರ್ಬಿರ್ಭು	₹	I	x	NHSO ₂ Q43	I	N-CH ₃	CN02	Z
1-110	a ₂ a ₂ a ₃ a	0420420H3	ĕ	I	I	x	ભૂવુ. ભૂતુ	ž	CCS	Ŧ
1-111	ರ್ಚಿದ್ಬು∞್ಡಿದ್ಯ	CH2CH2SO2CH3	퓽	∞CH ₃	I	I	Q42Q43	ž	CCOCH3	
1-112	G1200012013	CH2COCH2CH3	₹	OCH2CH3	I	I	۵	ž	C.SO ₃ H	Z
1-113	01201201	ठ ₂ ठभु	N. TOTA	I	I	NHCH3	ર્ફે_	Ž	CSO2NH2	2
1-114	1-114 G120000120F2	ಈ2ಯಂಗಿ2ಆ ₂ ಆ್2	I	ಯರಿಗ್ನರಗ್ರ	I	I	, o	ž	C.NH2	Zn
1-115	a12a12aN	0420420A	₹	I	x	QP ₂	£	¥	COCH	Z
1-116	CH2CH=CH2	CH2CH=CH2	ë	r	Ξ	Ŷ	ફ	ž	Ŷ	ន
1-117	012012013·	೦೫2೦೫2೦೫3	₹	x	=	ģ	ğ.	Ŧ	CCH2/C	Zu
1-118	Ŷ	I	8	I	I	r	ર્ફ	¥	C-NHCOCH ₃	ន
1-119	CH2CH3	ದ್ಬಚ್ಚ	Ж03н	Ι	I	NHSO ₃ H	Ŧ	¥	C-NHSO ₂ CH ₃	છે
1.120	012013	G-I ₂ Q-I ₃	жозн	Η.	r	\$02¢H₃	=	ž	C-N(CH ₃) ₂	2

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Compound	Ą	ፚ	ď	ď	ď	ď	45	×	>	Metal
1-121	1-121 Combined with R ₂	CH2 CH2 NHCOCH3	₹	Ξ	Ŧ	Ŧ	I	N-CH ₃	C-CH-CHCH3	Ž
1-122	1-122 Combined with R	Combined with R	₹	(CH ₂) ₃	x	(CH ₂) ₃	x	N-CH ₃	£	ž
1-123	1-123 Combined with R.	Combined with R.	퓽	an, wah,	=	CH2COCH2	=	¥	CSCN	Z
1-124	0420420H	4 ,44,04	₹	I	Ξ.	I	₹	N-CH ₃	೧.502	Z
1-125	a420420H	CH2CH2CH	8 H	r	r	I	I	NHN	C-CH*NCH2CH3 NI	Z .n
1-126	<u>ರಸ್ತಿದ್ದಾರ್</u> ಕ್ಕ	OH2CH2CH3	₹	x	æ	I	8	N-CH ₂	S	. ₹
1-127	ದ್ಯಚ್	ф ² Ф	₹	r	x	x	ર્જ	NCH2CH2CH3	r ccn	Z
1-128	CH2CH2CH2CH3	042012013	30 ₃ H	I	r	π.	3	NCH2CH2CH2 CCN	Pa CCN	Z

5		Metal	Ž	Z	2	Z	Z	ž	5	Z	Z	2	ភ	Z	Z	2	2
		>	z	Z	z	· Z	z	z	z	Z	z	z	z	z	z	z	z
		×	ဟ	Ø	ဟ	ဟ	တ	တ	Ø	တ	S	Ø	တ	ဟ	တ	ဟ	Ø
10		R,	\$\diamonds\$	ക്	₽	ă	I	I	r	0	B	HOO	r	CH2CH3	ರ್.	अभ्यत्र	ਨੰ
15 20		 Bu	×	90H3	яО ₃ н	I	x	I	Ö	r	<u> </u>	. ₹	NHCOCH,	I	I	x	I
		G _R	Ŧ	r	I	x	I	I	I	I	I	I	I	I	x	I	=
25		cz*	π	x	H ₂	I	포	x	σ	x	I	I	x	. エ	I	I	x
30		R	₹	NHOOCH	NHCOCH3	NHCOCH ₃	₹	₹	NHCOCH ₃	₹	₹	NHOO-H	NHCOCH ₃	₹	₽	Б	ቼ
35			ર્દ	ਨੂੰ	^	CH(CH ₃) 2		Ŧ.	H ₃)2	ź	ź		£		ગુવ્યક	ร์	ž
40	1)	B	ಯ್ಯಯ _ೆ ಯ್ಯಯ್ಯ	CH2CH2CCH2CH3	\$ 25 E	GH2CH2CH2C	CH2CH3	ದ್ಯು ದ್ಬುದ್ಬು	CH2CH(CH3)2	ದ್ಯು ದ್ಯುದ್ಯು	⇔.⇔.⇔.⇔.	CH2CH2CH	കുകു കുക	CH2CH3	CH2CH2NHSO2CH3	<u> </u>	⇔₂⇔₂⇔₃⇔₃
45 50	Table 1 (continued)	F.	ರ್ಚಿದ್ಯಾರ್ಡ್ಕ	ಈ ₂ ಯ್2ಯ,	1-131 042	1-132 04204204(043)2 042042004204(043)2	ಈ 2 ಈ 3	1-134 CH2CH2CH3	1-135 CH2CH(CH3)2	1-136 CH2CH2CH3	1-137 042042043	1-138 сн2сн2сн	1-139 GH2CH2CCH2CH	ರ್ನೆಚ್	1-141 CH2CH2NHSO2CH3	⇔್ತದ್ಯ⇔,	1.143 G2Q2Q2Q3
55	Tab	Compound	1-129	1-130	1-131	1-1320	1-133	1-134	1-135 (1-136	1-137	1-138	1-139	1-140	1-141	1-142	1-143

Ta	Table 1 (continued)	1)								
Compound	ď.	ъ.	P3	2	q.	.a.	R,	×	>	Metal
1.144	Q12Q13	Cl ₂ Gl ₃	HOCO	Ξ	ェ	¥	20 20	S	z	7
1-145	G1200H2CH20CH3	0420CH20CH3	1 000	I	I	x	కో	ဟ	z	Z
1-146	CH2 CH2 NHCOCH3	CH2CH2NHCOCH3	8 H	H000	I	I	8	ဟ	z	Z
1-147	9129420042CH(CH3)2 9429420H(CH3)2	, 04204200H2CH(CH3)2	80 ₃ H	80 ₃ H	I	803H	NO2	တ	z	Z
1-148	Q4,2Q43	ದ್ದ ಚ್ಚ	NHSO2Q43	NH3	I	r	8	ဟ	ž	Ž
1-149	G12012012013	ರ ₂ ರ್2ರ್ಚರು	₹	I	x	NHSO ₂ CH ₃	ж 2 сн ³	ဟ	z	Z
1-150	042042013	ರ್ಚಿಚ್ಚಾರ್ಚ್ಯ	ŏ	x	I	x	ಯ ,ಯಯ,ಯ	ဟ	z	Z
1-151	G42G42SO2G43	a124250243	8	80 F	I	x	CHNaNCH2CH3	ဟ	z	్ట్రీ
1-152	04200042043	CH2COCH2CH3	₹	OCH ₂ GH ₃	I	Έ-	N(CH(CH ₃) ₂) ₂	ဟ	z	Z
1-153	CH2CH2CH	GH2 GH2 GH	NHCOCH	r	I	NHCH ₃	80 H	တ	z	Pd
1-154	CH2COCCH2CF2CHF2	ದ₁ ಯಂ ೧ಗ₁೧೯₂೧4೯₂	ĕ	COCH2CH3	I	I	SO ₃ H	တ	z	Zu Zu
1-155	क ृक्रु	Q2Q2Q	₽	I	I	P Š	NH3	တ	z	Z
1.156	CH2CH=CH2	CH2CH=CH2	ĕ	I	I	0	0	ဟ	z	ន
1-157	⇔₂⇔₂⇔₃	<u>ದ್ಯಾರ್</u> ಚ್ಚರ್ಚು	₹	Ι	Ξ	ą,	000H20H3	S	z	vz.
1-158		I	803H	I	I	Ξ	∞0014 ₃	ω .	z	Zu

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Compound	æ	œ ^z	ڇ	æ æ	α _ε ,	ھے	ት	×	>	X Y Metal
1-159	ದ್ಬಿದ್ಯ	d ₁ d ₁	80 ₃ H	r	Ξ	HEOSH	MHSO ₃ H CH ₂	Ø	z	්රී
1-160	ದ್ಯುದ್ಯ	ଫଃଫ	SO ₃ H	I	x	∞_2 CH3	NHCOCI43 S	S	2	Z
1-161	Combined with R, CH2CH2NHCOCH3	CH2CH2NHCOCH3	용	I	x	I	NHSO ₂ CH ₃ S	တ	z	Ž
1-162	Combined with R, Combined with R, OH	Combined with R	₹_	(CH ₂) ₃	Ĭ	(CH ₂) ₃	CH=CHCH3 S	တ	z	Z
1-163	1-163 Combined with R, Combined with R, OH	Combined with R	₹_	CH2COCH2	I	CH2COCH2	х	ဟ	z	Z

Table 1 (continued)

[0060] In the present invention, a phthalocyanine compound contained in the dye layer is the phthalocyanine compound represented by Formula (2) having an absorption maximum in a wavelength range of 680 to 900 nm and having preferably a large refractive index and a small absorbance in a wavelength range of 770 to 830 nm.

[0061] When the dye layer comprises a two-layer structure of the recording layer and the optical interference layer, the optical constants (refractive index n and attenuation coefficient k) required for the recording layer containing the phthalocyanine compound are 1.8 or more for n and 0.04 to 0.20 for k at the wavelength λ_1 of the laser bearns, and 1.1 or more for n and 0.04 to 0.6 for k at the wavelength λ_2 .

[0062] If n is a smaller value than the above range at the wavelength λ_1 , it is difficult to obtain the reflectance and the modulation degree which satisfy the CD standards. The value of n smaller than the above range also at the wavelength λ_2 makes it difficult to obtain the reflectance needed for reading signals accurately.

[0063] Meanwhile, the value of k exceeding 0.20 at the wavelength λ_1 lowers the reflectance and makes it difficult to satisfy the CD standards, and the value of k less than 0.04 makes recording impossible. The value of k exceeding 0.6 at the wavelength λ_2 not only increases the absorbance too much and does not provide the reflectance needed for reproduction but also makes the signals to be liable to change by reproducing light. Accordingly, it is not suited to practical use. Considering recording at the wavelength λ_2 , k has to be 0.04 or more.

[0064] In the substituents Y₁ to Y₈ in Formula (2) described above, the hydrocarbon group includes a saturated hydrocarbon group such as methyl, ethyl, butyl, pentyl, hexyl, heptyl, octyl, nonyl, decyl, dodecyl, cyclohexyl and dimethylcyclohexyl, and unsaturated hydrocarbon group such as ethenyl, butenyl, hexenyl, octenyl, dodecenyl, butynyl, heptynyl, phenyl, methylphenyl, butylphenyl and hexylphenyl. These hydrocarbon groups may be linear, branched or cyclic and may be substituted with halogen, an amino group, a cyano group, an ether group or hydroxyl group.

[0065] A halogen atom includes fluorine, chlorine, bromine and iodine.

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[0066] The substitution positions of the substituents of Y_1 to Y_6 and A_1 to A_4 bonded to the benzene rings constituting a phthalocyanine ring shall not specifically be restricted, and the kind and number of the substituents may be different for the four benzene rings in the molecule.

[0067] The divalent metal atom represented by M includes Cu, Zn, Fe, Co, Ni, Ru, Pd, Pt, Mg, Ti, Be, Ca, Ba, Pb and Cd, and one-substituted trivalent metal includes Al-Cl, Al-Br, Ga-Cl, Ga-Br, In-Cl, In-Br, Ti-Cl, Ti-Br, Al-C₆H₅, Al-C₆H₄(CH₃), In-C₁₀H₇, Mn(OH), Mn(OC₆H₅), Mn(OSi(CH₃)₃), FeCl and RuCl, and two-substituted tetravalent metal includes CrCl₂, SiCl₂, CeBr₂, SnCl₂, TiCl₂, Mn(OH)₂, Sn(OH)₂, TiR₂, CrR₂, SiR₂, SnR₂, GeR₂ (R represents an alkyl group, a naphthyl group or a derivative thereof), Ti(OR')₂, Cr(OR')₂, Si(OR')₂ and Sn(OR')₂ (R' represents an alkyl group, a phenyl group, a naphthyl group, a trialkylsityl group, a dialkylalkoxysityl group or a derivative thereof), and oxy metal includes VO, MnO and TiO.

[0068] The concrete examples of the above phthalocyanine compound include the following compounds (Formulas (a) to (c)). To be detailed, given are the compounds described in Japanese Patent Application Laid-Open No. 3-62878 (1991), Japanese Patent Application Laid-Open No. 3-141582 (1991) and Japanese Patent Application Laid-Open No. 3-215466 (1991), and they can be synthesized by the methods described in these publications.

In Formula (a) described above, the compounds having the substituents and metals shown below:

Compound	М	Α	R
a-1	Pd	Н	OCH(CH ₃)CH(CH ₃) ₂
a-2	Pd	н	OCH[CH(CH ₃) ₂] ₂
a-3	Pd	н	OCH[CH(CH ₃)(C ₂ H ₅)][CH(CH ₃) ₂]
a-4	Pd	Н	OCH ₂ CH(C ₂ H ₅)C ₄ H ₉
a-5	Ni	NO ₂	OCH[CH(CH ₃) ₂] ₂
a-6	Ni	н	S-n-C ₈ H ₁₇
a-7	Ni	н	S(CH ₂ CH ₂ O) ₂ CH ₃
a-8	Cu	н	OCH[CH(CH ₃)(C ₂ H ₅)][CH(CH ₃) ₂]
a-9	VO	н	OCH ₂ CH(C ₂ H ₅)C ₄ H ₉
a-10	Cu	н	OCH ₂ CH ₂ OCH ₂ CH ₂ OCH ₃
a-11	vo	Н	OCH ₂ (CF ₂) ₃ CF ₂ H

In Formula (b) described above, the compounds having the substituents and metals shown below:

Compound	М	Χn	P.
b-1	Pd	Br _{2.5}	OCH(CH ₃)CH(CH ₃) ₂
b-2	Pd	Cl ₈	OCH[CH(CH ₃) ₂] ₂
b-3	Pd	Br ₄	OCH[CH(CH ₃)(C ₂ H ₅)][CH(CH ₃) ₂]
b-4	Pd	Br ₄	OCH(CH ₃)CH(CH ₃) ₂
b-5	Pd	14	OCH(CH ₃)CH(CH ₃) ₂
b-6	Pd	13.2	OCH ₂ CH(C ₂ H ₅)C ₄ H ₉
b-7	Pđ	Br ₄	OCH[CH(CH ₃) ₂] ₂
b-8	Pd	Br ₄	OCH(CH3)CH2CH(CH3)2
b-9	Pd	Br _{3.5}	OCH(CH ₃)CH(CH ₃) ₂

In Formula (c) described above, the compounds having the substituents and metals shown below:

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Compound	М	A'	R"	P**
c-1	Pd	Н	O(CH ₂) ₂ CH(CH ₃) ₂	O(CH ₂) ₂ CH(CH ₃) ₂
c-2	Pd	CI	O(CH ₂) ₂ CH(CH ₃) ₂	O(CH ₂) ₂ CH(CH ₃) ₂
c-3	Pd	н	O-n-C ₅ H ₁₁	O-n-C ₅ H ₁₁
c-4	Pd	CI	O(CH ₂) ₂ CH(CH ₃) ₂	O-n-C ₄ H ₉
c-5	Pd	Н	OCH2CH(C2H5)C4H9	OCH ₂ CH(C ₂ H ₅)C ₄ H ₉
c-6	Ni	SPh	O-n-C ₈ H ₁₇	O-n-C ₈ H ₁₇
c-7	Cu	CI	O(CH ₂) ₂ CH(CH ₃) ₂	O(CH ₂) ₂ CH(CH ₃) ₂
c-8	Al	CI	O(CH ₂) ₂ CH(CH ₃) ₂	O(CH ₂) ₂ CH(CH ₃) ₂
c-9	vo	CI	O-n-C ₈ H ₁₇	O-n-C ₈ H ₁₇
c-10	Cu	CI	O-n-C ₈ H ₁₇	O-n-C ₈ H ₁₇
c-11	Cu	CI	O(CH ₂) ₂ N(C ₂ H ₅) ₂	OCH3
c-12	Ni	Н	O-n-C ₈ H ₁₇	O-n-C ₈ H ₁₇

[0069] These phthalocyanine compounds may be used alone or in a mixture of the plural compounds.

[0970] A method for providing the dye layer in the present invention includes, for example, dipping, spraying, spin coating, casting, sputtering, chemical deposition and vacuum deposition, and the spin coating is preferred since it is simple.

[0071] When a coating method such as spin coating is used, a coating solution prepared by dissolving or dispersing the azo compound represented by Formula (1) or the metal complex thereof and/or the phthalocyanine compound represented by Formula (2) (hereinafter referred to as the dye) in a range of 0.05 to 30 weight %, preferably 0.5 to 20 weight % is used. In this case, a solvent which does not damage the substrate is preferably selected. The solvent includes, for example, alcohol series solvents such as methanol, ethanol, isopropanol, octafluoropentanol, allyl alcohol, methyl cellosolve, ethyl cellosolve and tetrafluoropropanol, aliphatic or alicyclic hydrocarbon series solvents such as hexane, heptane, octane, decane, cyclohexane, methylcyclohexane, ethylcyclohexane and dimethylcyclohexane, aromatic hydrocarbon series solvents such as toluene, xylene and benzene, halogenated hydrocarbon solvents such as carbon tetrachloride, chloroform, tetrachloroethane and dibromoethane, ether solvents such as diethyl ether, dibutyl ether, diisopropyl ether and dioxane, ketone solvents such as acetone and 3-hydroxy-3-methyl-2-butanone, ester solvents such as ethyl acetate and methyl lactate, and water. They may be used alone or in a mixture of plural solvents. [0072] When the dye layer comprises a two-layer structure of the recording layer and the optical interference layer, the optical interference layer each dissolved in different polar solvents are used, and each is preferably not dissolved in the optical interference layer each dissolved in different polar solvents are used, and each is preferably not dissolved in different polar solvents are used, and each is preferably not dissolved.

in the other solvent.

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[0073] The azo compound or the metal complex (dye) thereof used in the optical interference layer is dissolved in high polar solvents, for example, alcohol solvents and water, and coated, and the phthalocyanine compound (dye) used in the recording layer is dissolved in a low polar solvent, for example, aliphatic or alicyclic hydrocarbon solvents, aromatic hydrocarbon solvents, carbon tetrachloride and ether solvents, and coated. In order to decrease damage to a resin substrate or influence to a multilayer film, solvents for coating and forming the respective dye layers are preferably an aliphatic or allcyclic hydrocarbon solvent and a mixed solvent thereof with other solvents, and an alcohol solvent and a mixed solvent thereof with other solvents is particularly desired.

10 [0074] It may be possible, if necessary, to disperse the dyes for the dye layer into a polymer film.

[0075] When solvents which do not damage the substrate can not be selected, sputtering, chemical deposition and vacuum deposition are effective.

[0076] A film thickness of the dye layer shall not specifically be restricted and is preferably 50 to 300 nm. The film thickness of the dye layer smaller than 50 nm not only makes recording impossible or causes distortion on recording signals because of large heat diffusion, but also decreases signal amplitudes. As a result the CD standards are not satisfied. Meanwhile, the film thickness larger than 300 nm lowers the reflectance to deteriorate the reproducing signal characteristics.

[0077] When the dye layer comprises a two-layer structure of the recording layer and the optical interference layer, an average film thickness of the optical interference layer satisfies the following equation at the wavelengths of λ_1 and λ_2 and preferably is a thickness at which the reflectance becomes larger at the respective laser wavelengths:

70≤ni x di≤300

- wherein ni is a refractive index of the optical interference layer, and di is a film thickness of the optical interference layer. [0078] In the case where ni \times di is less than 70, the reflectance to light having the wavelength λ_2 is less than 15%, and the modulation degree decreases as well. In the case where ni \times di exceeds 300, the reflectance to light having the wavelength λ_1 is less than 65%.
- [0079] If necessary, additives such as a quencher, a thermal decomposition promotor of dyes, a UV absorber and an adhesive may be blended with the dyes described above, or groups having such effects may be introduced as substituents into the dyes.
 - [0080] Preferred as the quencher are metal complexes based on acetyl acetonates, bisdithio- α -diketones, bisdithiols such as bisphenyldithiol, thiocatechols, salicylaldehyde oximes and thiobisphenolates. Further, amines are suitable as well.
- 35 [0081] The thermal decomposition promoter shall not specifically be restricted as long as promotion of thermal decomposition of the dyes can be confirmed by means of thermogravimetric analysis (TG analysis), and includes metal compounds such as metalic antiknocking agents, metallocene compounds and acetylacetonato metal complexes.
 - [0082] Further, binders, leveling agents, and defoaming agents may be used as well in combination, if necessary. The preferred binder includes polyvinyl alcohol, polyvinyl pyrrolidone, nittrocellulose, cellulose acetate, ketone resins, acrylic resins, polystyrene resins, urethane resins, polyvinyl butyral, polycarbonate, polyolefins, and the like.
 - [0083] Further, dyes other than those described above may be added as well in order to improve recording characteristics.
 - [0084] The examples of the dyes include pentamethinecyanine dyes, heptamethinecyanine dyes, squalillium dyes, naphthoquinone dyes, azo dyes, naphthalocyanine dyes, phthalocyanine dyes and anthraquinone dyes. A blending ratio of these dyes falls in a level of 0.1 to 10 %.
 - [0085] In forming the dye layer on the substrate, a layer comprising inorganic substances or polymers may be provided on the substrate in order to improve the solvent resistance of the substrate, reflectance and the recording sensitivity.
- [0086] The content of the azo compound represented by Formula (1) or the metal complex thereof and/or the phthalocyanine compound represented by Formula (2) in the dye layer is 30 % or more, preferably 60 % or more. It may be substantially 100 %.
 - [0087] Next, a reflective layer having a thickness of preferably 50 to 300 nm is formed on the dye layer described above. Substances providing sufficiently high reflectances at wavelengths of reproducing lights, for example, metals such as Au, Al, Ag, Cu, Ti, Cr, Ni, Pt, Ta, Cr and Pd can be used as a material for the reflective layer alone or in the form of alloys. Among them, Au and Al have the high reflectances and therefore are suitable to the material for the reflective layer. In addition to the above substances, the following ones may be contained. There may be included, for example, metals and semi-metals such as Mg, Se, Hf, V, Nb, Ru, W, Mn, Re, Fe, Co, Rh, Ir, Cu, Zn, Cd, Ga, In, Si, Ge, Te, Pb, Po, Sn and Bi. A substance containing Au as a main component is suitable since the reflective layer having

a high reflectance can readily be obtained. The term "main component" means a component having a content of 50 % or more. A multilayer film may be formed for the reflective layer with materials other than metals by laminating a thin film having a low refractive index after the other.

[0088] Methods for forming the reflective layer include, for example, sputtering, ion plating, chemical deposition and vacuum deposition. Further, a known inorganic or organic intermediate layer and an adhesive layer may be provided on the substrate or under the reflective layer in order to improve the reflectance and the recording characteristics.

[0089] Further, a protective layer is formed on the reflective layer in order to protect the dye layer and the reflective layer. Two sheets of the media may be stuck together.

[0090] Materials for the protective layer shall not specifically be restricted as long as they can protect the reflective layer from external force. The organic materials include thermoplastic resins, thermosetting resins, electron beam-curing resins, and UV-curing resins. The inorganic materials include SiO₂, SiN₄, MgF₂, and SnO₂. The protective layer can be formed by applying a coating solution prepared by dissolving a thermoplastic resin or a thermosetting resin in a suitable solvent, and drying it. A UV curing resin is applied as it is or as a solution prepared by dissolving it in a suitable solvent, and it is irradiated with UV rays to cure, whereby the protective layer can be formed. Acrylate resins such as, for example, urethane acrylates, epoxy acrylates and polyester acrylates can be used as the UV-curing resin. These materials may be used alone or in a mixture, and they may be used not only in a single layer film but also in a multilayer film.

[0091] Coating methods such as spin coating and casting, and methods such as sputtering and chemical deposition are used as a method for forming the protective layer as is the case with the recording layer. Among them, the spin coating is preferred.

[0092] In general, the film thickness of the protective layer falls in a range of 0.1 to 100 μ m. In the present invention, however, it is 3 to 30 μ m, preferably 5 to 20 μ m.

[0093] Further, printing such as labeling can be applied on the protective layer.

[0094] A film of a UV-curing resin or an inorganic film may be formed on a specular side of the substrate in order to protect the surface and prevent dusts from sticking thereon.

[0095] The optical recording medium of the present invention thus obtained can record and reproduce by focusing laser beams on the dye layer. Signals used in recording include, for example, EMF modulation signals used for CD. The medium of the present invention can be recorded and reproduced with laser beams having a wavelength of about 780 nm since the reflectance of 65 % or more to light having a wavelength selected from 770 to 830 nm can be obtained. Further, recorded information can be reproduced with commercial CD and CD-ROM players. The characteristics of the signals reproduced satisfy the orange book standards which are the standards of CD-R. Further, the medium of the present invention provides a reflectance of 15 % or more to light having a wavelength falling in a range of 620 to 690 nm and therefore can be reproduced with an optical disc player for high density recording media loaded with a laser having a wavelength selected from arrange of 620 to 690 nm. Light used for players for high density recording media in the next age has a wavelength falling in a range of 620 to 690 nm. Lasers having wavelengths falling in this range include dye lasers, which have wavelengths to be selected in a wide range of a visible ray region, and a helium-neon laser having a wavelength of 633 nm. Further, semiconductor lasers which are put into practical use have a wavelength of, for example, 635 nm, 650 nm or about 680 nm. The medium of the present invention can be recorded as well with light having a wavelength selected from a range of 620 to 690 nm.

EXAMPLES

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[0096] The examples of the present invention shall be shown below but the present invention shall by no means be restricted by these examples.

Example 1:

[0097] The azo compound (1-19) of 0.2 g shown in Table-1 out of the azo compounds represented by Formula (1) and the phthalocyanine compound of 0.02 g represented by the chemical formula (a-11) described previously were dissolved in diacetone alcohol (produced by Tokyo Kasei K.K.) of 10 ml to prepare a dye solution. Used as a substrate was a disc (made of polycarbonate resin) having a diameter of 120 mm and a thickness of 1.2 mm with a spiral pregroove (track pitch: 1.6 µm).

[0098] The dye solution was spin-coated on this substrate at a revolution of 1500 rpm and dried at 70°C for 2 hours to form a recording layer. Au was sputtered on this recording layer with a sputtering equipment (CDI-900: manufactured by Balzers Co., Ltd.) to form a reflective layer having a thickness of 100 nm. Sputtering was carried out at a gas pressure of 1.33 Pa (1.0 x 10°2 Torr).

[0099] Further, a UV-curing resin SD-17 (manufactured by Dainippon Ink Chemical Ind. Co., Ltd.) was spin-coated on the reflective layer and then irradiated with UV rays to form a protective layer having a thickness of 6 µm.

[0100] Recording was carried out on the optical recording medium thus obtained at a linear velocity of 5.6 m/s and a laser power of 10 mW with an optical disc evaluation apparatus DDU-1000 (manufactured by Pulstec Ind. Co., Ltd.) loaded with a red semiconductor laser head having a wavelength of 680 nm, and an EFM encoder (manufactured by Kenwood Co., Ltd.). After recording, the signals were reproduced with an evaluation apparatus loaded with a red semiconductor laser head having a wavelength of 635 nm to determine the reflectance.

[0101] Further, this recorded sample was reproduced and evaluated with a commercial CD player having a reproducing wavelength of 780 nm to find that the sample showed good recording characteristics.

Examples 2 to 28:

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[0102] Optical recording media were produced in the same manner as described in Example 1, except that the azo compounds described in Table-1 and the phthalocyanine compounds represented by the chemical formulas (a-10 to 11) described previously were used in combination.

[0103] Recording was carried out on the optical recording media thus obtained in the same manner as that in Example 1 at a linear velocity of 5.6 m/s and a laser power of 10 mW with the optical disc evaluation apparatus DDU-1000 (manufactured by Pulstec Ind. Co., Ltd.) loaded with the red semiconductor laser head having a wavelength of 680 nm and the EFM encoder (manufactured by Kenwood Co., Ltd.). After recording, the same measurements as those in Example 1 were carried out to find that the good recording characteristics were shown in any cases.

20 Example 29:

[0104] First, a solution prepared by dissolving 0.8 g of the phthalocyanine compound represented by the chemical formula (b-7) described previously in dimethylcyclohexane (produced by Tokyo Kasei K.K.) of 40 ml was used to form a recording layer by spin-coating. Then, the azo compound (1-31) described in Table-1 was used to form an interference layer on this recording layer on the same conditions as those in Example 1, whereby an optical recording medium comprising two layers of the interference layer and the recording layer was prepared.

[0105] Recording was carried out on the optical recording medium thus produced at a linear velocity of 2.8 m/s and a laser power of 8 mW with the optical disc evaluation apparatus DDU-1000 (manufactured by Pulstec Ind. Co., Ltd.) loaded with a near infrared semiconductor laser head having a wavelength of 780 nm, and the EFM encoder (manufactured by Kenwood Co., Ltd.). After recording, the same measurements as in Example 1 were carried out to find that the good recording characteristics were shown in any cases.

[0106] Further, the signals were reproduced with an evaluation apparatus loaded with a red semiconductor laser head having a wavelength of 680 nm to find that such good recording characteristics as the reflectance of 31 % were shown.

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Examples 30 to 46:

[0107] Optical recording media were produced in the same manner as that described in Example 29, except that the azo compounds described in Table-1 and the phthalocyanine compounds represented by the chemical formulas (b-1 to c-10) described previously were used in combination.

[0108] Recording was carried out on the optical recording media thus produced at a linear velocity of 2.8 m/s and a laser power of 8 mW with the optical disc evaluation apparatus DDU-1000 (manufactured by Pulstec Ind. Co., Ltd.) loaded with the near infrared semiconductor laser head having a wavelength of 780 nm, and the EFM encoder (manufactured by Kenwood Co., Ltd.). After recording, the same measurements as those in Example 1 were carried out to find that the good recording characteristics were shown in any cases.

[0109] Further, the signals were reproduced with the evaluation apparatus loaded with the red semiconductor laser head having a wavelength of 680 nm to find that such good recording characteristics as the reflectances of 30 % or more were shown in the respective cases.

50 Example 47:

[0110] First, a solution prepared by dissolving 0,8 g of the phthalocyanine compound represented by the chemical formula (b-7) described previously in dimethylocyclohexane (produced by Tokyo Kasei K.K.) of 40 ml was used to form a recording layer by spin-coating. Then, the azo compound (1-83) described in Table-1 was used to form an interference layer on this recording layer on the same conditions as those in Example 1, whereby an optical recording medium comprising two layers of the interference layer and the recording layer was prepared.

[0111] Recording was carried out on the optical recording medium thus produced at a linear velocity of 5.6 m/s and a laser power of 10 mW with the optical disc evaluation apparatus DDU-1000 (manufactured by Pulstec Ind. Co., Ltd.)



loaded with the red semiconductor laser head having the wavelength of 680 nm, and the EFM encoder (manufactured by Kenwood Co., Ltd.). After recording, the same measurements as those in Example 1 were carried out to find that the good recording characteristics were shown in any cases.

[0112] Further, the signals were reproduced with the evaluation apparatus loaded with the red semiconductor laser head having a wavelength of 680 nm to find that such good recording characteristics as the reflectance of 30 % were shown.

Example 48:

[0113] An optical recording medium was produced in the same manner as that described in Example 47, except that the azo compound (1-39) described in Table-1 and the phthalocyanine compound (b-7) represented by the chemical formula described previously were used in combination.

[0114] Recording was carried out on the optical recording medium thus produced at a linear velocity of 5.6 m/s and a laser power of 10 mW with the optical disc evaluation apparatus DDU-1000 (manufactured by Pulstec Ind. Co., Ltd.) loaded with the red semiconductor laser head having a wavelength of 680 nm, and the EFM encoder (manufactured by Kenwood Co., Ltd.). After recording, the same measurements as those in Example 1 were carried out to find that the good recording characteristics were shown in any cases.

[0115] Further, the signals were reproduced with the evaluation apparatus loaded with the red semiconductor laser head having a wavelength of 680 nm to find that such good recording characteristics as the reflectance of 30 % were shown.

[0116] Shown in Table 2 together are the measured results of the reflectances in reproducing with the red semiconductor laser having a wavelength of 635 nm and the reflectances and the error rates in reproducing with a commercial CD player.

Table 2

	Chen	nical formula No.		Reproduction	· · · · · · · · · · · · · · · · · · ·
Example	Azo	Phthalocyanine	635 nm	780	nm
			Reflectance (%)	Reflectance (%)	Error rate (cps)
1	1-19	a-11	27.9	69.5	6
2	1-20	a-11	31.5	71.2	7
3	1-21	a-11	29.4	70.6	5
4	1-22	a-11	29.6	71.4	8
5	1-23	a-11	31.2	69.3	5
6	1-24	a-11	30.1	70.5	9
7	1-25	a-10	34.3	71.8	5
8	1-26	a-10	33.2	70.4	5
9	1-27	a-10	32.7	73.1	7
10	1-28	a-10	32.1	72.3	4
11	1-29	a-10	31.4	71.0	6
12	1-30	a-10	33.8	69.1	7
13	1-63	a-11	34.0	72.3	6
14	1-64	a-11	31.2	72.4	6
15	1-65	a-11	31.6	73.4	4
16	1-66	a-11	34.7	71.5	6
17	1-67	a-11	32.1	73.2	8
18	1-68	a-11	30.4	71.4	5
19	1-69	a-11	31.3	72.1	4
20	1-70	a-11	33.3	71.3	5
21	1-71	a-10	29.8	73.1	6
22	1-72	a-10	33.2	69.7	7
23	1-73	a-10	33.0	71.6	7
24	1-74	a-10	29.7	70.0	9
25	1-75	a-10	28.3	69.2	5



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Table 2 (continued)

	Chem	nical formula No.		Reproduction	
Example	Azo	Phthalocyanine	635 nm	780	nm .
			Reflectance (%)	Reflectance (%)	Error rate (cps)
26	1-76	a-10	32.7	70.9	7
27	1-77	a-10	32.7	70.7	8
28	1-78	a-10	31.5	71.3	5
29	1-31	b-7	32.6	71.3	4
30	1-32	b-7	33.0	70.9	5
31	1-34	c-10	29.6	72.8	4
32	1-35	c-9	32.2	71.0	7
33	1-36	c -7	29.3	69.7	. 8
34	1-37	b-8	32.3	72.2	8
35	1-38	b-1	28.8	70.7	6
36	1-30	b-6	31.6	71.5	7
37	1-79	b-7	33.3	73.7	4
38	1-80	b-7	33.4	72.8	4
39	1-81	b-7	32.1	70.5	5
40	1-82	b-7	27.6	73.9	8
41	1-84	c-10	29.7	71.8	7
42	1-85	c-9	31.2	71.0	8
43	1-86	c-7	30.6	72.7	8
44	1-87	b-8	28.4	68.6	6
45	1-88	b-1	30.0	70.3	6
46	1-78	b-6	31.2	72.3	5
47	1-83	b-7	31.4	72.1	5
48	1-39	b-7	32.1	70.3	6

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Example 49:

- 35 [0117] The metal complex (1-93) of the azo compound of 0.2 g shown in Table-1 out of the metal complexes of the azo compounds represented by Formula (1) and 0.02 g of the phthalocyanine compound (a-10) represented by the chemical formula described previously were dissolved in 2,2,3,3-tetrafluoro-1-propanol of 10 ml to prepare a dye solution. Used as a substrate was a disc having a diameter of 120 mm and a thickness of 1.2 mm, which was made of polycarbonate resin and had a spiral pre-groove (track pitch: 1.6 μm).
 - [0118] The dye solution was spin-coated on this substrate at a revolution of 1500 rpm and dried at 70°C for 2 hours to form a recording layer. Au was sputtered on this recording layer with the sputtering equipment (CDI-900: manufactured by Balzers Co., Ltd.) to form a reflective layer having a thickness of 100 nm. Sputtering was carried out at a gas pressure of 1.33 Pa (1.0 x 10°2 Torr). Further, the UV-curing resin SD-17 (manufactured by Dainippon Ink Chemical Ind. Co., Ltd.) was spin-coated on the reflective layer and then irradiated with UV rays to form a protective layer having a thickness of 6 µm.
 - [0119] EFM signals were recorded on this optical recording medium at a linear velocity of 2.8 m/s and a laser power of 8 mW with a writer (CDD-521: manufactured by Philiphs Co., Ltd.). After recording, the signals were reproduced with a commercial CD player (YAMAHA CDX-1050; laser wavelength: 786 nm) to determine the reflectance, the error rate and the modulation degree. As a result, the distortion of the reproducing wave form was small, and therefore the good values satisfying the orange book standards were shown. Next, the signals recorded on this medium were reproduced with the optical disc evaluation apparatus (DDU-1000: manufactured by Pulstec Ind. Co., Ltd.) loaded with a red semiconductor laser head having wavelengths of 680 nm and 635 nm to determine the reflectance, the error rate and the modulation degree (I3/Itop) of the shortest pit. It has been confirmed that the good values are shown in any cases.
 - [0120] Next, recording was carried out on this medium at a linear velocity of 5.6 m/s and a laser power of 10 mW with the optical disc evaluation apparatus (DDU-1000: manufactured by Pulstec Ind. Co., Ltd.) loaded with the red semiconductor laser head having a wavelength of 680 nm, and the EFM encoder (manufactured by Kenwood Co.,

Ltd.). After recording, the signals were reproduced with an evaluation apparatus loaded with a red semiconductor laser head having wavelengths of 680 nm and 635 nm to determine the reflectance, the error rate and the I3/Itop. The good values were shown in any cases. The signals recorded on the medium with the above drive (DDU-1000) having a wavelength of 680 nm were reproduced with the commercial CD player (YAMAHA CDX-1050; laser wavelength: 786 nm) to determine the reflectance, the error rate and the I3/Itop. The good values satisfying the orange book standards were shown in any cases.

Examples 50 to 54:

10 [0121] Optical recording media were produced in the same manner as that described in Example 49, except that the metal complexes of the azo compounds described in Table-1 out of the metal complexes of the azo compounds represented by Formula (1), and the phthalocyanine compounds represented by the chemical formulas (a-10 to 11) described previously were used in combination (shown in Table 3).

[0122] The media thus produced recorded and was evaluated in the same manners as those in Example 49. As a result, it has been confirmed that the good values are shown in any cases.

Example 55:

[0123] The metal complex (1-89) of the azo compound of 0.2 g shown in Table-1 out of the metal complexes of the azo compounds represented by Formula (1) was dissolved in 2,2,3,3-tetrafluoro-1-propanol of 10 ml to prepare a dye solution 1.

[0124] Used as a substrate was a disc having a diameter of 120 mm and a thickness of 1.2 mm, which was made of polycarbonate resin and had a spiral pre-groove (170 nm thick, 0.5 µm wide, track pitch: 1.6 µm).

[0125] The dye solution 1 was spin-coated on this substrate at a revolution of 1500 rpm and dried at 70°C for 3 hours to form a dye optical interference layer. With respect to the optical constants of this optical interference layer, the refractive index was 1.9 and the attenuation coefficient was 0.05 at 780 nm; the refractive index was 2.1 and the attenuation coefficient was 0.04 at 680 nm; and the refractive index was 2.5 and the attenuation coefficient was 0.10 at 635 nm. Accordingly, the products (ni × di) of the refractive indices and the film thickness at 780, 680 and 635 nm were 152, 168 and 200, respectively.

[0126] Further, a dye solution 2 prepared by dissolving 0.25 g of the phthalocyanine compound (c-2) represented by the chemical formula described previously in 1,2-dimethylcyclohexane of 10 ml was spin-coated at a revolution of 1600 rpm and dried at 70°C for 2 hours to form a recording layer. With respect to the optical constants of this recording layer, the refractive index was 2.2 and the attenuation coefficient was 0.08 at 780 nm; the refractive index was 1.2 and the attenuation coefficient was 0.34 at 635 nm.

[0127] Next, Au was sputtered on this recording layer with the sputtering equipment (CDI-900: manufactured by Balzers Co., Ltd.) to form a reflective layer having a thickness of 100 µm. Further, the UV-curing resin SD-17 (manufactured by Dainippon Ink Chemical Ind. Co., Ltd.) was spin-coated on the reflective layer, and then irradiated with UV rays to form a protective layer having a thickness of 6 µm, whereby an optical recording medium was prepared.

[0128] EFM signals were recorded on this optical recording medium at a linear velocity of 2.8 m/s and a laser power of 8 mW with the writer (CDD-521: manufactured by Philiphs Co., Ltd.). After recording, the signals were reproduced with the commercial CD player (YAMAHA CDX-1050; laser wavelength: 786 nm) to determine the reflectance, the error rate and the modulation degree. As a result, the distortion of the reproducing wave form was small, and therefore the good values satisfying the orange book standards were shown. Next, the signals recorded on this medium were reproduced with the optical disc evaluation apparatus (DDU-1000: manufactured by Pulstec Ind. Co., Ltd.) loaded with the red semiconductor laser head having wavelengths of 680 nm and 635 nm to determine the reflectance, the error rate and the modulation degree (I3/Itop) of the shortest pit. The good values were shown in any cases.

[0129] Next, recording was carried out on this medium at a linear velocity of 5.6 m/s and a laser power of 10 mW with the optical disc evaluation apparatus (DDU-1000: manufactured by Pulstec Ind. Co., Ltd.) loaded with the red semiconductor laser head having a wavelength of 680 nm and the EFM encoder (manufactured by Kenwood Co., Ltd.). After recording, the signals were reproduced with the evaluation apparatus loaded with the red semiconductor laser head having wavelengths of 680 nm and 635 nm to determine the reflectance, the error rate and the I3/Itop. The good values were shown in any cases. The signals recorded on the medium with the above drive (DDU-1000) having a wavelength of 680 nm were reproduced with the commercial CD player (YAMAHA CDX-1050; laser wavelength: 786 nm) to determine the reflectance, the error rate and the I3/Itop. The good values satisfying the orange book standards were shown in any cases.

Examples 56 to 58:

[0130] Optical recording media each having a recording layer formed on an optical interference layer were produced in the same manner as that described in Example 55, except that the metal complexes of the azo compounds described in Table-1 out of the metal complexes of the azo compounds represented by Formula (1), and the phthalocyanine compounds represented by the chemical formulas (b-3 and c-2) described previously were used in combination (shown in Table-3).

[0131] The media thus produced was recorded at 780 nm and 680 nm and reproduced at 786 nm, 680 nm and 635 nm in the same manners as those in Example 55. As a result, the good recording characteristics were shown in any of the above wavelengths.

Example 59:

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[0132] An optical recording medium was prepared in the same manner as that described in Example 55, except that the coating order was reversed and that the phthalocyanine compound (c-2) represented by the chemical formula described previously was applied for a recording layer and the metal complex (1-89) of the azo compound described in Table-1 was applied thereon for an optical interference layer.

[0133] The medium thus prepared was recorded and reproduced in the same manners as those in Example 55. As a result, the good recording characteristics were shown in any of the wavelengths.

Examples 60 to 133:

[0134] Optical recording media each having an optical interference layer formed on a recording layer were prepared and evaluated in the same manners as those described in Example 59, except that the metal complexes of the azo compounds described in Table-1 and the phthalocyanine compounds represented by the chemical formulas (a-6 to c-2) described previously were used in combination.

[0135] Described in Table-3 are the combinations of the dyes, the optical constants [refractive index (n) and attenuation coefficient (k)] of the recording layers and the optical interference layers at 780, 680 and 635 nm, the film thicknesses of the optical interference layers and the ni × di values.

30 [0136] All the media had large modulation degrees and small error rates and jitters in a recording mode of high to low at the respective wavelengths of 780, 680 and 635 nm, and therefore very good reproductions could be made.



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		49	50	51	52	53	54	55	56	57	58	59	9	63	62
Recording layer formula No	rmula No.	8-10	8-11	a-10	a-10	a-11	a-10	C-2	b-3	2-5	p-3	c-2	b-3	G-2	D-0
		1-93.	1-94	1-94	1-133		1-134 1-134								
Interference layer formula No.	formula No	•						68-1	1-90	1-129	1-130	1-89	1-90	1-89 1-90 1-129 1-130 1-89 1-90 1-129 1-130	1-130
film th	film thickness (nm)	~						80	100	80	100	75	100	75	100
Optical constants															
Recording layer	780 nm n°1							2.2	2.3	2.2	2.3	2.2	2.2	2.2	2.2
		_						0.08	0.12	0.08	0.12	0.08	0.08	0.08	90.0
	680 nm n							1.2	1.3	1.2	1.3	1.2	1.2	1.2	1.2
	×							0.49	0.51	0.49	0.51	0.49	0.49	0.49	0.49
	635 nm n							1.2	1.2	1.2	1.2	1.2	1.2	1.2	1.2
	×							0.34	0.30	0.34	0.30	0.34	0.34	0.34	0.34
Interference	780 nm n							1.9	1.8	1.9	1.8	1.9	1.9	1.9	1.9
layer	×						Ĭ	0.05	90.0	0.05	90.0	0.12	0.05	0.12	0.05
	680 nm n							2.1	1.9	2.1	1.9	2.1	2.1	2.1	2.1
	*						_	0.04	0.05	p.04	0.05	0.12	0.04	0.12	0.04
	635 nm n							2.5	2.2 . 2.4	2.4	2.2	5.6	2.5	5.6	2.5
	×							0.10	0.10	0.10	0.10	0.14	0.10	0.14	0.10
nt × d.t.	780 חש							152	180	152	180	143	190	143	190
	680 nm							168	190	168	190	158	210	158	210
	635 nm							200	220	200	220	195	250	195	253

*1 n: refractive index *2 k: attenuation coefficient *3 refractive index x film thickness in interference layer

Table 3 (continued)

								Еха	Example				
			63	63 64 65	4	65	99	67	99	69	70	71	72
Recording layer f	formula No.	No.	a-11	.1 a-11		a-11	a-10	a-10	a-10	a-11	a-11	a-11	a-11
Interference layer formula No.	forn	nula N	lo. 1-91		1-92 1	1-93	1-94	1-94 1-95	1-96	1-97	1-98	1-99	1-100
film thickness (nm)	tcknes	nu) ss	1) 80	06 (90	80	75	100	09	45	80	06
Optical constants													
Recording layer	780 nm	ה שנ	2.3		2.2 2	2.1	2.2	2.2 2.2	2.2	2.2	2.3	2.1	2.2
		* *	0.10	00.	0.11.0	0.08	0.10	0.09	0.09	0.11	0.13	0.13 0.08 0.07	0.07
	680 r	n n	1.2	1.3		1.3	1.2	1.2 1.2	1.2	1.2	1.2	1.2	1.3
		×	0.48	8 0.50		0.48	0.46	0.48	0.49	0.45	0.46	0.49	0.43
	635 r	n En	1.2		1.1	1.2	1.1	1.2	1.1 1.2 1.2	1.1	1.1	1.2	1.2
		×	0.34	4 0.30		0.31	0.29	0.29 0.32	0.34	0.29	0.27	0.31	0.30
Interference	780 n	r Ec	1.9	1.8		1.9	1.9	1.9	1.9	1.9	1.8	1.9	1.9
layer		×	0.05	5 0.10		10	0.10 0.11	0.12	0.05	0.11	0.12	0.05 0.10	0.10
	680 n	C EC	2.1	1.9		0	5.0	2.1	2.0 2.0 2.1 2.1 2.0	2.0	2.0	2.0 2.1 2.0	2.0
		×	0.04	4 0.08		10	0.10	0.10 0.10 0.12	0.04 0.10	0.10	0.11	0.04	0.10
	635 n	ת שת	2.5	2.4		2.4	2.5	2.6 2.5		2.5	5.4	2.5	2.4
		×	0.10	0 0.11		0.12	0.13	0.14	0.10	0.13	0.15	0.10	0.12
ni x di*³	780 n	mc1	152	162		171	152	143	190	114	81	152	171
	680 n	E	168	171		180	160	158	210	120	90	168	180
	635 n	E C	200	216		216	200	195	250	150	108	200	216

*1 n: refractive index *2 k: attenuation coefficient

Table 3 (continued)

T3 74 75 76 77 78 79 80 Cmula No.							Example	ple				
. c-2 c-2 c-2 c-2 b-3 b-3 b-3 b-3 b-3 hondown) 90 90 90 80 75 100 60 45 hondown) 90 90 90 80 75 100 60 45 hondown) 90 90 90 80 75 100 60 45 hondown) 90 90 90 80 75 100 60 45 hondown) 90 90 90 80 75 100 60 45 hondown 1.2 1.2 2.1 2.0 2.1 2.2 kg 0.07 0.11 0.09 0.07 0.11 0.10 hondown 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.3 hondown 1.1 1.2 1.3 1.2 1.1 1.2 1.3 hondown 1.1 1.2 1.1 1.2 1.3 hondown 1.1 1.2 1.1 1.2 1.1 1.2 1.3 hondown 1.1 1.2 1.3 1.2 1.3 hondown 1.1 1.2 1.3 1.3 hondown 1.1 1.2 1.3 hondown 1.1 1.3 hondown 1			73	74	75	76	77	78	79	90	81	82
No. 1-101 1-102 1-103 1-104 1-105 1-106 1-107 1-108 nm) 90 90 90 80 75 100 60 45 n** 2.1 2.2 2.1 2.2 2.1 2.0 2.1 2.2 k** 0.07 0.11 0.07 0.11 0.09 0.07 0.11 0.10 n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.1 1.2 1.3 k 0.46 0.50 0.48 0.46 0.47 0.48 0.44 0.46 n 1.1 1.2 1.1 1.2 1.1 1.2 1.2 1.1 1.2 1.3 k 0.31 0.31 0.29 0.29 0.33 0.34 0.27 0.28 n 1.9 1.8 1.9 1.9 1.9 1.9 1.9 1.9 1.9 c 0.10 0.10 0.10 0.11 0.12 0.05 0.11 0.12 n 2.0 1.9 2.0 2.0 2.1 2.1 2.0 2.0 c 0.10 0.08 0.10 0.10 0.12 0.05 0.11 0.12 n 2.4 2.4 2.4 2.5 2.6 2.5 2.5 2.4 c 0.12 0.11 0.12 0.13 0.14 0.10 0.13 0.15 171 162 171 152 143 190 114 81 180 171 180 160 158 210 120 90	Recording layer fo	ormula No.	c-2		0-2	c-2	b-3	p-3	p-3	p-3	p-9	6-q
780 nm n°1 2.1 2.2 2.1 2.2 2.1 2.0 2.1 680 nm n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 780 nm n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.1 1.2 685 nm n 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 780 nm n 1.9 1.8 1.9 1.9 1.9 1.9 1.9 1.9 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.1 2.0 685 nm n 2.0 1.0 0.18 0.10 0.12 0.13 0.14 0.10 0.13 780 nm 171 162 171 180 160 158 210 120 785 250 150	Interference layer	formula			1-103	1-104		1-106	1-107	1-108	1-109	1-110
780 nm n°1 2.1 2.2 2.1 2.2 2.1 2.0 2.1 680 nm n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.1 1.2 1.3 1.2 1.1 1.1	film thi	tckness (n		06	90	80	75	100	90	45	80	90
780 nm n°1 2.1 2.2 2.1 2.2 2.1 2.0 2.1 680 nm n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.1 1.2 1.3 1.2 1.1 1.2 1.1 1.2 1.3 1.2 1.1 1.2 1.2	Optical constants											
680 nm n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.1 1.2 680 nm n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.2	Recording layer		1 2.1		2.1	2.2	2.1	2.0	2.1	2.2	2.1	2.0
680 nm n 1.2 1.3 1.2 1.3 1.2 1.1 1.2 1.1 1.2 635 nm n 1.1 1.2 1.2					0.07	0.11	0.09	0.07	0.11	0.10	0.07	0.02
635 nm n 1.1 1.2 1.1 1.2 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.1 1.2 1.2			1.2		1.2	1.3	1.2	1.1	1.2	1.3	1.2	1.3
635 nm n 1.1 1.2 1.1 1.2 1.2 1.1 1.2 780 nm n 1.9 1.8 1.9 1.9 1.9 1.9 1.9 1.9 680 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.0 635 nm n 2.4 2.4 2.4 2.5 2.6 2.5 2.5 780 nm n 1.9 1.1 0.12 0.13 0.14 0.10 0.13 780 nm n 1.0 1.1 1.1 1.1 1.1 1.1 1.1 1.1 1.1 1.1			0.4		0.48	0.46	0.47	0.48	0.44	0.46	0.48	0.46
K 0.31 0.31 0.29 0.29 0.33 0.34 0.27 780 nm 1.9 1.9 1.9 1.9 1.9 1.9 1.9 680 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.0 635 nm n 2.4 2.4 2.4 2.4 2.5 2.6 2.5 2.5 780 nm 171 162 171 180 160 158 210 0.13 680 nm 180 171 180 160 158 210 120 680 nm 180 171 180 160 158 210 120			1.1		1.1	1.2	1.2	1.1	1.2	1.2	1.1	1.2
780 nm n 1.9 1.8 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9		×	0.3		0.29	0.29	0.33	0.34	0.27	0.28	0.33	0.30
680 nm n 2.0 1.9 2.0 2.0 2.1 2.1 2.0 6.11 k 0.10 0.00 0.10 0.12 0.05 0.11 k 0.10 0.00 0.10 0.10 0.12 0.04 0.10 6.35 nm n 2.4 2.4 2.4 2.5 2.6 2.5 2.5 780 nm 171 162 171 152 143 190 114 6.10 0.13 0.14 0.10 0.15 0.14 0.10 0.15 0.15 0.15 0.10 0.15 0.15 0.15	Interference		1.9		1.9	1.9	1.9	1.9	1.9	1.8	1.9	1.9
680 nm n 2.0 1.9 2.0 2.0 2.1, 2.1 2.0 k 0.10 0.00 0.10 0.12 0.04 0.10 635 nm n 2.4 2.4 2.4 2.5 2.6 2.5 2.5 2.5 k 0.12 0.11 0.12 0.13 0.14 0.10 0.13 780 nm 171 162 171 152 143 190 114 680 nm 180 171 180 160 158 210 120 0.15 2.15 2.15 2.15 2.15 2.15 2.15 2.15 2	laver		0.1		0.10	0.11	0.12	0.05	0.11	0.12	0.05	0.10
635 nm n 2.4 2.4 2.4 2.5 2.6 2.5 2.5 2.5 2.5 2.5 2.5 2.5 2.5 2.5 2.5			2.0		2.0	5.0	2.1	2.1	5.0	5.0	2.1	2.0
635 nm n 2.4 2.4 2.4 2.5 2.6 2.5 2.5 2.5 2.5 2.5 2.5 2.5 2.5 2.5 2.5			0.1		0.10	0.10	0.12	0.04	0.10	0.11	0.04	0.10
780 nm 171 162 171 152 143 190 114 680 nm 180 171 180 160 158 210 120			2.4		2.4	2.5	5.6	2.5	2.5	2.4	2.5	2.4
780 nm 171 162 171 152 143 190 114 680 nm 180 171 180 160 158 210 120			0.1		0.12	0.13	0.14	0.10	0.13	0.15	0.10	0.12
680 nm 180 171 180 160 158 210 120	nt x dt		171		171	152	143	190	114	81	152	171
250 150		mu 089	180		180	160	158	210	120	90	168	180
25 C7 C7 C17 C17 C17		635 nm	216		216	200	195	250	150	108	200	216

*1 n: refractive index *2 k: attenuation coefficient *3 refractive index x film thickness in interference layer

Table 3 (continued)

							Example	ple				
			83	84	85	98	87	88	89	90	91	92
Recording layer for	er formula No	No.	c-2	c-2	c-2	c-2	p-3	b-3	p-3	p-3	p-9	p-9
	layer formula No.	la No.	1-111	1-112	1-113	1-114	1-115	1-116	1-117	1-118	1-119	1-120
	lm thickness	(ma)	80	90		80	75	100	90	54	80	90
Optical constants												
Recording layer	780 nm		2.1	2.3	2.2	2.1	2.2	2.1	2.2	2.3	2.1	2.1
		<u>.</u> *	0.11	0.11	0.07	60.0	0.08	0.09	0.11	0.10	0.08	0.07
	680 nm	c	1.2	1.3	1.2	1.3	1.3	1.2	1.2	1.1	1.2	1.3
		×	0.46	0.50	0.46	0.46	0.50	0.48	0.45	0.46	0.48	0.46
	635 nm	c	1.2	1.1	1.1	1.2	1.1	1.2	1.2	1.2	1.1	1.2
		×	0.29	0.31	0.30	0.29	0.32	0.33	0.29		0.33	0.29
Interference	780 nm	c	1.9	1.8	1.9	1.9	1.9	1.9	1.9		1.9	1.9
laver		×	0.11	0.10	0.10	0.11	0.12	0.05	0.11		0.05	0.10
	680 nm	c	2.0	1.9	2.0	2.0	2.1	2.1	5.0	2.0	2.1	5.0
		×	0.10	0.08	0.10	0.10	0.12	0.04	0.10	0.11	0.04	0.10
	635 nm	c	2.4	2.4	2.4	2.5	5.6	2.5	2.5	2.4	2.5	2.4
		×	0.12	0.11	0.12	0.13	0.14	0.10	0.13	0.15	0.10	0.12
nt × d1.	780 nm		152	162	171	152	143	190	114	81	152	171
	680 nm		160	171	180	160	158	210	120	90	168	180
	635 nm		200	216	216	200	195	250	150	108	200	216

*1 n: refractive index *2 k: attenuation coefficient *3 refractive index x film thickness in interference layer

Table 3 (continued)

Recording layer formula No. Interference layer formula No. film thickness (nm) Optical constants Recording layer 780 nm n k************************************	93 a = B	č	L	90	07	ao	00	טטו
Recording layer formula No. Interference layer formula film thickness (1) Optical constants Recording layer 780 nm 1		Į,	75	2		22		777
Interference layer formula film thickness (1 Optical constants Recording layer 780 nm 1		a-8	a-8	p-2	p-2	b-2	p-2	p-2
, id	No. 1	1-122	1-123	1-124	1-125	1-126 1-127	1-127	1-128
780 nm 780 m 680 m .	•	06		80	75	100	09	45
mir 089		2,2	2.0	2.2	2.1	2.2	2.2	2.3
Ē	., 2.1 k** 0.08		0.07	0.11	0.08	0.09	0.11	0.09
			1.3	1.2	1.3	1.2	1.3	1.2
•	0.48	0.50	0.47	0.46	0.49	0.48	0.45	0.46
635 nm	1.2	1.1	1.2	1.2	1.1	1.2	1.1	1.2
	0.29	0.30	0.31	0.29	0.34	0.33	0.28	0.28
780 nm 780 nm	1.9	1.8	1.9	1.9	1.9	1.9	1.9	1.8
	0.17	0.11	0.10	0.11	0.12	0.08	0.10	0.11
19787 TO	2.0	1.9	2.0	2.0	2.1	2.1	5.0	5.0
	0.10	0.09	0.11	0.10	0.12	0.06	0.12	0.11
ma 26.35	2.4	2.4	2.4	2.5	2.6	2.5	2.5	2.4
	0.12	0.11	0.12	0.13	0.13	0.11	0.12	0.14
780 pm	171	162	171	152	143	190	114	81
680	180	171	180	160	158	210	120	90
ma 253	216	216	216	200	195	250	150	108

*1 n: refractive index *2 k: attenuation coefficient *3 refractive index x film thickness in interference layer

Table 3 (continued)

							Ехап	Ехамріе				
		İ	101	102	103	104	105	106	107	108	109	110
Recording layer fo	yer formula No	o.	p-3	p-3	p-3	b-3	p-3	p-9	p-9	6-q	p-9	6-q
	layer formula No.	a No.	1-131	1-132	1-133	1-134	1-135	1-136	1-137 1-138	1-138	1-139	1-140
	ilm thickness (nm)	(ma)	90	06	90	80	75	100	90	45	80	90
Optical constants												
Recording layer	780 nm	r c	2.2	2.3	5.0	2.2	2.2	2.2	2.5	2.3	2.2	2.0
		*	0.08	0.12	90.0	0.10	0.08	0.08	0.10	0.11	0.08	90.0
	mu 089	c	1.2	1.3	1.3	1.2	1.2	1.2	1.2	1.2	1.2	1.3
		×	0.49	0.51	0.47	0.45	0.49	0.49	0.45	0.45	0.49	0.47
	635 nm	c	1.2	1.2	1.2	1.1	1.2	1.2	1.1	1.1	1.2	1.2
		×	0.34	0.30	0.30	0.28	0.34	0.34	0.28	0.27	0.34	0.30
Interference	780 nm	c	1.9	1.8		1.9	1.9	1.9		1.8	1.9	1.9
		×	0.05	0.10	0.10	0.11	0.12	0.05		0.12	0.05	0.10
	680 nm	c	2.1	1.9	2.0	2.0	2.1	2.1	5.0	5.0	2.1	2.0
		×	0.04	0.08	0.10	0.10	0.12	0.04	0.10	0.11	0.04	0.10
	635 nm	c	2.5	2.4	2.4	2.5	2.6	2.5	2.5	2.4	2.5	2.4
		×	0.10	0.11	0.12	0.13	0.14	0.10	0.13	0.15	0.10	0.12
ni x di"	780 nm		152	162	171	152	143	190	114	81	152	171
			168	171	180	160	158	210	120	06	168	180
	635 nm	,	200	216	216	200	195	250	150	108	200	216

*In: refractive index *2 k: attenuation coefficient *3 refractive index x film thickness in interference layer

Table 3 (continued)

						Exa	Example		! 		
		111	112	113	114	7.	116	117			
Recording layer formula, No.	formula No.	p-9	p- 0		A-6	9-6		,	110	777	120
Interference lav	Layer formula Me			•	•		0-5	a-0	a-6	a-6	a-6
	C. TOTHILLA NO.	1-141	1-142	1-143	1-144	1-145	1-146	1-147	1-148	_	
tilm	film thickness (nm)	90		0	a	ū					067-7
· Optical constants		•	2	2	3	Ç	007	09	45	80	90
Recording layer	r 780 nm n*1	,	c	•	•						
			5.3	2.0	2.2	7.5	2.2	2.2	2.3	2.2	2.0
		0.06	0.12	90.0	0.10	0.08	0.08	0.10	0.11	0.08	90
	u wu ORq	1.3	1.3	1.3	1.2	1.2	1.2	1.2	1.2	1 2) .
	×	0.47	0.51	0.47	0.45	0 40	0			3 .	2.1
	635 nm n	,	•					5	0.45	0.49	0.47
		7:1	7.7	1.2	1.1	1.2	1.2	1.1	1.1	1.2	1.2
		0.30	0.30	0.30	0.28	0.34	0.34	0.28	0.27	0.34	0
anna ranna	780 nm n	1.9	1.8	1.9	1.9	1.9		0		5 ,	
layer	*	01.0	0	5				h • •	0.1	۲.۶	1.9
	. m. 089		21.	0.10	77.0	0.12	0.05	0.11	0.12	0.05	0.10
		0.4	6.1	2.0	2.0	2.1		2.0	2.0	2.1	2.0
		0.10	0.08	0.10	0.10	0.12		0.10	0.11	0.04	0.10
	035 mm n	2.4	2.4	2.4	2.5	2.6		2.5	2.4	2.5	2 4
	×	0.12	0.11	0.12	0.13	0.14	0.10	0.13	0.15	2	; ;
ום א זח	780 nm	171	162	171	152	143		114	81.0		3
	680 nm	180	171	180	160	158	210	120		707	7/7
	635 nm	216	216	216				2	2	901	780
			1	640	2077	195	250	150	108	200	216

Table 3 (continued)

							Exar	Example				
			121	122	123	124	125	126	127	128	129	130
Recording layer formula No.	ormula	NO.	a-6	a-6	a-6	8-8	8-8	8-8	8-8	8-8	8-8	b-2
Interference layer	form	layer formula No.	1-151	1-152	1-153	1-154	1-155	1-156	1-157	1-158	1-159	1-160
film thickness (nm)	cknes	(mn) s	80	90	90	80	75	100	9	45	80	06
Optical constants											1	
Recording layer	780 nm		2.2	2.3	5.0	2.2	2.2	2.2	2.2	2.3	2.2	2.0
		<u>.</u> ×	0.10	0.12	0.06	0.10	0.08	0.08	0.10	0.11	0.08	90.0
	680 nm	C E	1.2	1.3	1.3	1.2	1.2	1.2	1.2	1.2	1.2	1.3
		×	0.45	0.51	0.47	0.45	0.49	0.49	0.45	0.45	0.49	0.47
	635 nm	E E	1.1	1.2	1.2	1.1	1.2	1.2	1.1	1.1	1.2	1.2
-		×	0.28	0:30	0:30	0.28	0.34	0.34	0.28	0.27	0.34	0.30
Interference	780 nm	ב	1.9	1.8	1.9	1.9	1.9	1.9	1.9	1.8	1.9	1.9
layer		×	0.11	0.10	0.10	0.11	0.12	0.05	0.11	0.12	0.05	0.10
	680 nm	ב	2.0	1.9	2.0	2.0	2.1	2.1	2.0	2.0	2.1	2.0
		×	0.10	0.08	0.10	0.10	0.12	0.04	0.10	0.11	0.04	0.10
	635 nm	נו	2.5	2.4	2.4	2.5	5.6	2.5	2.5	2.4	2.5	2.4
;		×	0.13	0.11	0.12	0.13	0.14	0.10	0.13	0.15	0.10	0.12
ni x di	780 nm		152	162	171	152	143	190	114	81	152	171
	mu 089		160	171	180	160	158	210	120	90	168	180
	635 nm		200	216	216	200	195	250	150	108	200	216

Comparative Example 1:

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[0137] An optical recording medium was prepared in the same manner as that in Example 55, except that the thickness of the optical interference layer comprising the azo compound was changed to 35 nm. The products (ni x di) of the refractive indices and the film thickness of the optical interference layer at 780, 680 and 635 nm were 67, 74 and 88, respectively.

[0138] The medium thus prepared was evaluated in the same manner as that described in Example 55 to find that the reproducing signal wave forms were distorted at the respective wavelengths of 680 and 635 nm, and the reflectance was low.

Comparative Example 2:

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[0139] An optical recording medium was prepared in the same manner as that described in Example 59, except that the thickness of the optical interference layer comprising the azo compound was changed to 160 nm. The products (ni \times di) of the refractive indices and the film thickness of the optical interference layer at 780, 680 and 635 nm were 304, 336 and 400, respectively.

[0140] The medium thus prepared was evaluated in the same manner as that in Example 59 to find that the reproducing signal wave forms were distorted at the respective wavelengths of 786, 680 and 635 nm, and the reflectance was low as well.

Comparative Example 3:

[0141] An optical recording medium was prepared in the same manner as that described in Example 49, except that only the metal complex (1-93) of the azo compound described in Table-1 was used in the dye layer.

[0142] The medium thus prepared was evaluated in the same manner as that in Example 49 to find that recording could not be carried out at 780 nm and the sensitivity was low at 680 nm, which therefore made fine recording impossible.

Comparative Example 4:

[01.43] An optical recording medium was prepared in the same manner as that described in Example 49, except that only the phthalocyanine compound (a-10) represented by the chemical formula described previously was used in the dye layer.

[0144] The medium thus prepared was evaluated in the same manner as that in Example 49 to find that the wave form was distorted and the reflectances at 635 and 680 nm were low.

[0145] Described in Table-4 together are the reproducing signal characteristics [reflectance (%), error rate (cps) and modulation degree (I3/Itop)] at 786, 680 and 635 nm when the respective media described above were recorded at 780 and 680 nm, respectively.

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	Example	64	50	15	53	53	2	1	1	1	
min 689	786 nm reproduction					2	,	66	20	72	28
recording	Reflectance (%)	99	68	70	70	69	70	69	67	69	99
	Error rate (cps)	6	7	7	æ	7	7	φ	S	, 4	\$ 5
	13/Itop	0.41	0.43	0.42	0.42	0.41	0.41	0.43	0.41	0.41	0.40
	680 nm reproduction										?
	Reflectance (%)	27	28	27	27	28	25	26	24	26	2.1
	Error rate (cps)	6	7	7	10	7	7	ס	. S	10	1 15
	13/Itop	0.38	0.40	0.39	0.38	0.40	0.39	0.42	0.40	0.42	0.40
	635 nm reproduction) !		?
	Reflectance (%)	33	31	32	33	31	32	32	32	32	32
	Error rate (cps)	7	9	9	7	Ŋ	Ŋ	o	7	10	2
	13/Itop	0.44	0.46	0.46	0.45	0.47	0.46	0.45	0.44	0.45	0.44
juu 082	786 nm reproduction) ;	; ;
recording	Reflectance (%)	20	69	89	71	68	29	69	70	69	20
	Error rate (cps)	9	7	9	9	7	9	10	9	10	, S.
	I3/Itop	0.45	0.43	0.44	0.45	0.43	0.44	0.44	0.43	0.44	0.43
	680 nm reproduction						_				}
	Reflectance (%)	28	. 42	27	28	27	. 27	30	29	30	29
	Error rate (cps)	Q	10	0	0	10	6	60	6	œ	6
	I3/Itop	0.39	0.46	0.41	0.39	0.46	0.41	0.38	0.45	0.38	0.45
	635 nm reproduction) -		
	Reflectance (%)	32	33	32	35	33	32	33	34	33	34
	Error rate (cps)	œ	6	9	80	6	7	7	<5	7	. 55
	I3/Itop	0.45	0.47	0.46	0.44	0.47	0.45	0.45	0.43	0.45	0.43

I3/Itop

(

5		65		68	9	0.40		26	œ	0.43		28	9	0.45		69	æ	0.47		30	7	0.42) -	31	α	,
10		64		70	~	0.45		28	O,	0.39		32	, 5	0.44		70	9	0.46		28	6	0.45		30	6	
		63		68	7	0.42		27	\$	0.46		30	\$	0.46		68	9	0.43		27	Ŋ	0.44		31	9	
15		62		70	\$	0.45		33	\$	0.45		28	7	0.43		99	< 5	0.47	•	28	80	0.41		33	\$	
20		61		69	, S	0.45		25	< 5	0.37		25	10	0.43		89	S	0.46		31	6	0.40		31	7	
		9		20		0.45		31	7	0.45		29	7	0.44		69	< 5	0.47		28	æ	0.41		33	< 5	
25		59		69	ထ	0.45		25	80	0.37		32	6	0.43		89	7	0.46		31	6	0.40		31	7	
30	ed)		uction	(%)	(cps)		uction	(%)	(cbs)		ction	(%)	cps)		ction	(%)	cps)		uction	(%)	(sdo		uction	(%)	Sps.)	
35	Table 4 (continued)	Ехамріе	786 nm reproduction	Reflectance (%)	Error rate (cps	13/Itop	680 nm reproduction	Reflectance (%)	Error rate (cps)	13/Itop	635 nm reproduction	Reflectance (%)	Error rate (cps)	I3/Itop	786 nm reproduction	Reflectance (%)	Error rate (cps)	13/Itop	680 nm reproduction	Reflectance (%)	Error rate (cps	I3/Itop	635 nm reproduction	Reflectance (%)	Error rate (cps.)	
40	Table		786	ď,	ជ		680	Ä.	ជ		635	Re	Er		786	Re	Er		680	Re	Er		635	Rei	Eri	
45			680 nm	recording											780 nm	recording										
50															•											

Table 4 (continued)

	Example	99	67	68	69	70	7.1	72	73	74	75
mr 089	786 nm reproduction										
recording	Reflectance (%)	70	69	70	69	70	69	20	68	69	69
	Error rate (cps)	7	60	9	7	9	\$	œ	9	\$	9
	I3/Itop	0.42	0.41	0.41	0.41	0.45	0.40	0.45	0.42	0.45	0.40
	680 nm reproduction										
	Reflectance (%)	27	28	27	30	26	28	31	26	28	28
	Error rate (cps)	σ	7	7	6	7	89	\$	80	10	80
	I3/Itop	0.39	0.41	0.39	0.44	0.38	0.38	0.43	0.45	0.41	0.42
	635 nm reproduction										
	Reflectance (%)	32	31	32	31	29	30	29	29	31	28
	Error rate (cps)	7	S	ĸ	80	01	Φ	7	တ	~	9
	13/Itop	0.46	0.45	0.44	0.45	0.46	0.47	0.43	0.45	0.44	0.45
. 180 பய்	786 nm reproduction										
recording	Reflectance (%)	70	69	68	89	69	71	99	69	20	69
	Error rate (cps)	9	7	7	6	9	Ŋ	σ	7	œ	80
	I3/Itop	0.44	0.43	0.44	0.44	0.45	0.43	0.47	0.43	0.46	0.47
	680 nm reproduction					_	_				
	Reflectance (%)	29	27	29	30	31	53	53	27	28	30
	Error rate (cps)	6	10	6	œ	0	6	œ	S	10	7
	I3/Itop	0.39	0.46	0.41	0.38	0.41	0.45	0.41	0.44	0.45	0.42
	635 nm reproduction										
	Reflectance (%)	34	32	31	34	32	34	33	31	30	31
	Error rate (cps)	æ	σ	80	7	7	< 5	< 5	9	٥,	ю
	I3/Itop	0.46	0.47	0.44	0.45 0.46	0.46	0.43	0.45	0.44	0.45	0.45

Table 4 (continued)

	Example	76	77	78	79	80	81	82	83	84	85
mr 089	786 nm reproduction	1)							
recording	Reflectance (%)	69	69	71	69	68	67	99	68	69	69
	Error rate (cps)	7	ω	9	7	9	Š	œ	9	4	9
	I3/Itop	0.41	0.42	0.41	0.42	0.41	0.42	0.43	0.42	0.44	0.40
	680 nm reproduction										
	Reflectance (%)	28	28	26	30	28	29	30	27	53	28
	Error rate (cps)	80	7	7	6	80	80	< 5	80	6	6
	I3/Itop	0.39	0.40	0.38	0.43	0.39	0.39	0.41	0.44	0.42	0.41
	635 nm reproduction										
	Reflectance (%)	32	31	31	31	28	30	28	29	30	28
	Error rate (cps)	7	ß	Ŋ	æ	6	80	7	Ŋ	\$	9
	I3/Itop	0.45	0.47	0.44	0.46	0.45	0.46	0.44	0.43	0.44	0.44
min 087	786 nm reproduction										
recording	Reflectance (%)	69	69	69	67	20	70	69	69	20	69
	Error rate (cps)	9	7	7	6	9	80	O,	7	80	60
	I3/Itop	0.43	0.43	0.45	0.46	0.45	0.44	0.43	0.42	0.43	0.44
	680 nm reproduction					-					
	Reflectance (%)	29	28	27	30	31	30	29	28	27	30
	Error rate (cps)	σ	80	6	ω	6	6	00	9	6	7
	I3/Itop	0.39	0.45	0.42	0.39	0.41	0.44	0.42	0.43	0.44	0.42
	635 nm reproduction										
	Reflectance (%)	34	33	32	31	32	33	33	31	30	31
	Error rate (cps)	80	6	80	7	7	<. 5	Š	9	Q	æ
	I3/Itop	0.45	0.46	0.45	0.44	0.46	0.43	0.46	0.45	0.46	0.45

Table 4 (continued)

	Example	86	87	88	89	90	91	92	93	94	95
680 nm	786 nm reproduction										
recording	Reflectance (%)	69	69	20	69	69	69	20	68	20	68
	Error rate (cps)	6	89	7	9	7	<5	ς>	7	\$	9
	I3/Itop	0.42	0.43	0.43	0.44	0.45	0.42	0.44	0.43	0.45	0.40
	680 nm reproduction										
	Reflectance (%)	28	28	27	26	27	25	30	27	28	29
	Error rate (cps)	6	80	2	0	8	\$	7	\$	6	80
	I3/Itop	0.39	0.41	0.42	0.41	0.39	0.40	0.43	0.44	0.40	0.42
	635 nm reproduction										
	Reflectance (%)	32	31	31	30	30	31	53	30	31	29
	Error rate (cps)	œ	œ	7	6	6	7	80	, 5	<. 5	9
	I3/Itop	0.44	0.45	0.46	0.42	0.43	0.43	0.44	0.44	0.44	0.45
780 nm	786 nm reproduction										
recording	Reflectance (%)	68	69	69	69	68	70	69	68	69	69
	Error rate (cps)	80	7	7	σ	7	9	۰ 5	7	9	80
	I3/Itop	0.43	0.44	0.45	0.44	0.43	0.43	0.45	0.42	0.44	0.44
	680 nm reproduction					_					
	Reflectance (%)	52	29	28	30	30	28	27	27	28	31
	Error rate (cps)	7	80	6	7	6	9	ω	S	0	7
	I3/Itop	0.39	0.40	0.42	0.39	0.41	0.43	0.41	0.42	0.41	0.42
	635 nm reproduction										
	Reflectance (%)	34	30	31	32	30	29	32	31	30	31
	Error rate (cps)	9	Ģ	ø	89	7	< 5	^ 2	9	O,	7
	13/Itop	0.44	0.43	0.46	0.45	0.46	0.43	0.45	0.44	0.45	0.45

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	Example	96	97	98	66	100	101	102	103
680 nm	786 nm reproduction								
recording	Reflactance (%)	68	69	20	69	69	29	69	99
	Error rate (cps)	6	7	7	9	ω	, 5	ŝ	9
	13/Itop	0.41	0.43	0.42	0.43	0.45	0.42	0.45	0.40
	680 nm reproduction								
	Reflectance (%)	27	28	28	26	25	26	28	21
	Error rate (cps)	9	7	7	0	89	< 5	10	œ
	I3/Itop	0.38	0.40	0.39	0.42	0.37	0.46	0.39	0.43
	635 nm reproduction								
	Reflectance (%)	31	32	31	32	32	30	32	28
	Error rate (cps)	7	9	9	O	6	<5	~	9
	I3/Itop	0.44	0.45	0.46	0.45	0.43	0.46	0.44	0.45
mr 082	786 nm reproduction								
recording	Reflectance (%)	69	69	68	69	68	68	70	69
	Error rate (cps)	• •	7	9	61	7	9	9	œ
	I3/Itop	0.44	0.43	0.44	0.44	0.46	0.43	0.46	0.47
	680 nm reproduction					-			
	Reflectance (%)	53	27	28	30	31	27	28	30
	Error rate (cps)	6	6	O,	ຍ	6	Ŋ	10	7
	13/1top	0.39	0.44	0.41	0.39	0.49	0.44	0.45	0.42
	635 nm reproduction								
	Reflectance (%)	34	32	32	33	31	31	30	31
	Error rate (cps)	6	O)	9	7	7	\$	9	10
	13/Itop	0.44	0.46	0.46	0.45	0.46	0.44	0.46	0.45

Table 4 (continued)

	Example	104	105	106	107	108	109	110	111	112	113
680 nm	786 nm reproduction										
recording	Reflectance (%)	71	69	71	69	20	69	70	68	69	69
	Error rate (cps)	ĸ	89	9	9	\$	\$	\$	9	<5	9
	I3/Itop	0.42	0.41	0.41	0.41	0.45	0.40	0.45	0.42	0.45	0.40
	680 nm reproduction										
	Reflectance (%)	27	28	25	30	25	21	31	26	28	23
	Error rate (cps)	10	7	7	10	7	89	\$	S	10	æ
	I3/Itop	0.38	0.40	0.39	0.42	0.37	0.40	0.45	0.46	0.39	0.43
	635 nm reproduction										
	Reflectance (%)	33	31	32	31	22	31	28	29	32	28
	Error rate (cps)	7	ß	S	10	O.	80	7	Ŋ	4	9
	I3/Itop	0.45	0.47	0.46	0.45	0.43	0.44	0.43	0.46	0.44	0.45
780 nm	786 nm reproduction										
recording .	Reflectance (%)	73	68	67	69	68	20	. 65	68	20	69
	Error rate (cps)	9	2	7	10	ø	S	10	7	9	80
	13/Itop	0.44	0.43	0.44	0.44	0.45	0.43	0.47	0.43	0.46	0.47
	680 nm reproduction					-					
	Reflectance (%)	53	27	29	30	31	29	29	27	28	30
	Error rate (cps)	6	10	01	ω	6	0	œ	ທ	10	7
	13/Itop	0.39	0.46	0.41	0.38	0.41	0.45	0.41	0.44	0.45	0.42
	635 nm reproduction										
	Reflectance (%)	35	33	31	33	31	34	33	31	30	31
	Error rate (cps)	æ	6	80	7	7	۸ ک	\$	9	10	80
	13/Itop	0.44	0.47	0.44	0.45	0.45	0.43	0.45	0.44	0.46	0.45

Claims

1. An optical recording medium comprising at least a dye layer, a reflective layer and a protective layer provided on a substrate, wherein an azo compound represented by the following Formula (1) or a metal complex thereof having an absorption maximum at a wavelength falling in a range of 450 to 630 nm and a phthalocyanine compound represented by the following Formula (2) having an absorption maximum at a wavelength falling in a range of 680 to 900 nm are contained in said dye layer.

$$R_7 \xrightarrow{Y-N} N=N \xrightarrow{R_3} R_4 R_1$$

$$R_5 \xrightarrow{R_6} R_6$$

wherein R_1 and R_2 represent independently a hydrogen atom, an alkyl group having 1 to 15 carbon atoms, an aryl group having 6 to 21 carbon atoms, an aralkyl group having 7 to 22 carbon atoms, or an alkenyl group having 2 to 16 carbon atoms;

 ${\rm R_3}$, ${\rm R_4}$, ${\rm R_5}$ and ${\rm R_6}$ represent independently a hydrogen atom, a halogen atom, a hydroxyl group, a carboxyl group, a sulfonic acid group, a sulfonamide group, an amino group, an alkyl group having 1 to 15 carbon atoms, an alkyl group having 1 to 15 carbon atoms, an aryl group having 6 to 21 carbon atoms, an acyl group having 1 to 15 carbon atoms, an alkylcarboxyl group having 2 to 16 carbon atoms, an aralkyl group having 2 to 22 carbon atoms, an alkylcarbonylamino group having 2 to 16 carbon atoms, an alkylsulfoamino group having 1 to 15 carbon atoms, an alkylsulfoamino group having 1 to 15 carbon atoms, or an alkenyl group having 2 to 16 carbon atoms.

 R_1 and R_4 , R_2 and R_6 , and R_1 and R_2 may form rings; R_7 represents a hydrogen atom, a halogen atom, a hydroxyl group, a carboxyl group, a sulfonic acid group, a sulfonamide group, an amino group, an alkyl group having 1 to 15 carbon atoms, an aryl group having 6 to 21 carbon atoms, an acyl group having 1 to 15 carbon atoms, an alkylcarboxyl group having 2 to 16 carbon atoms, an aralkyl group having 7 to 22 carbon atoms, an alkylcarboxylamino group having 2 to 16 carbon atoms, an alkylsulfoamino group having 1 to 15 carbon atoms, an alkylsulfoamino group having 1 to 15 carbon atoms, an alkylsulfoamino group having 1 to 15 carbon atoms, an alkylsulfonyl group having 1 to 15 carbon atoms, an alkylsulfonyl group having 1 to 15 carbon atoms, an alkylsulfonyl group having 3 to 16 carbon atoms, an alkylsulfonyl group, a mercapto group, a thiocyano group, a chlorosulfonic acid group, an alkylthio group having 1 to 15 carbon atoms, an alkylsulfonyl group having 2 to 16 carbon atoms, or an alkylsulfonyl group having 1 to 15 carbon atoms; X represents a sulfur atom or N-R₈, wherein R₈ represents a hydrogen atom, an alkyl group having 1 to 15 carbon atoms, an aryl group having 6 to 21 carbon atoms, an aralkyl group having 7 to 22 carbon atoms, or an alkenyl group having 2 to 16 carbon atoms; Y represents a nitrogen atom or C-R₉, wherein R₈ is synonymous with those described in R₇; provided that when X is a sulfur atom, Y is a nitrogen atom, and when X is N-R₈, Y is C-R₉:

$$(A_{4})_{m4} \qquad (Y_{2})_{m} \qquad (A_{1})_{m1} \qquad (Y_{3})_{14} \qquad (Y_{4})_{12} \qquad (Z)$$

$$(A_{4})_{m4} \qquad (Y_{5})_{14} \qquad (Y_{5})_{14} \qquad (Y_{5})_{12} \qquad (Z)$$

$$(A_{3})_{m3} \qquad (Y_{5})_{23} \qquad (Z)$$

wherein Y_1 , Y_2 , Y_3 , Y_4 , Y_5 , Y_6 , Y_7 , and Y_8 represent independently a hydrogen atom, a hydrocarbon group having

1 to 20 carbon atoms, an alkoxyl group having 1 to 20 carbon atoms, or an alkylthlo group having 1 to 20 carbon atoms; in the respective combinations of Y_1 and Y_2 , Y_3 and Y_4 , Y_5 and Y_8 and Y_7 and Y_8 , they may be combined to form rings when they are adjacent to each other; A_1 , A_2 , A_3 and A_4 represent independently a halogen atom or a nitro group; I_1 , I_2 , I_3 and I_4 represent an integer of 0 to 3; m_1 , m_2 , m_3 and m_4 represent an integer of 0 to 3; and M represent two hydrogen atoms, a divalent metal atom, a trivalent or tetravalent substituted metal atom, or an oxy metal.

The optical recording medium as claimed in claim 1, wherein the compound contained in the dye layer and having an absorption maximum at a wavelength falling in a range of 450 to 630 nm is a metal complex of an azo compound represented by the following Formula (3):

wherein R_1 , R_2 , R_4 , R_5 , and R_6 are synonymous with those described in Formula (1); R_{10} represents a hydroxyl group or a carboxyl group; and R_{11} represents a hydrogen atom or a halogen atom.

- 3. The optical recording medium as claimed in claim 1, wherein the dye layer comprises a single layer structure containing a mixture of the azo compound represented by Formula (1) or the metal complex thereof having an absorption maximum at a wavelength falling in a range of 450 to 630 nm and the phthalocyanine compound represented by Formula (2) having an absorption maximum at a wavelength falling in a range of 680 to 900 nm.
- 4. The optical recording medium as claimed in claim 1, wherein the dye layer comprises a two-layer structure of an optical interference layer containing the azo compound represented by Formula (1) or the metal complex thereof having an absorption maximum at a wavelength falling in a range of 450 to 630 nm and a recording layer containing the phthalocyanine compound represented by Formula (2) having an absorption maximum at a wavelength falling in a range of 680 to 900 nm.
- 5. The optical recording medium as claimed in claim 4, wherein an equation of 70≤ni x di≤300 applies to light -having a wavelength used for recording and reproducing, wherein ni represents a real part of a complex refractive index and di represents a layer thickness in the optical interference layer.
- 6. The optical recording medium as claimed in any of claims 1 to 5, capable of recording and/or reproducing with a laser beam having a wavelength λ₁ selected from wavelengths falling in a range of 770 to 830 nm and capable of recording and/or reproducing as well with a laser beam having a wavelength λ₂ selected from wavelengths falling in a range of 620 to 690 nm.
- The optical recording medium as claimed in claim 6, capable of reproducing with a laser beam having the wavelength λ₂.
 - 8. The optical recording medium as claimed in claim 6 or 7, wherein the laser beam having the wavelength λ_1 has a reflectance of 65 % or more, and the laser beam having the wavelength λ_2 has a reflectance of 15 % or more, which are measured through the substrate.
 - 9. The optical recording medium as claimed in claim 8, wherein the laser beam having the wavelength λ_2 has a reflectance of 20 % or more, which is measured through the substrate.
- A metal complex of an azo compound represented by Formula (4).

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$$\begin{bmatrix} R_{11} & N - N \\ S & N = N - N \end{bmatrix} = N \begin{bmatrix} R_1 \\ R_2 \end{bmatrix} M e^{2t}$$
 (4)

wherein R₁ and R₂ represent independently a hydrogen atom, an alkyl group having 1 to 15 carbon atoms, an aryl group having 6 to 21 carbon atoms, an aralkyl group having 7 to 22 carbon atoms, or an alkenyl group 2 to 16 carbon atoms; R₁₁ represents a hydrogen atom or a halogen atom; Me represents nickel, cobalt, copper, palladium, iron and zinc.

Patentansprüche

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1. Optisches Aufzeichnungsmedium, umfassend zumindest eine Farbschicht, eine reflektierende Schicht und eine auf einem Substrat angeordnete Schutzschicht, worin eine Azoverbindung der folgenden Formel (1) oder ein Metallkomplex hiervon mit einem Absorptionsmaximum bei einer Wellenlänge im Bereich von 450 bis 630 nm und eine Phthalocyaninverbindung der folgenden Formel (2) mit einem Absorptionsmaximum bei einer Wellenlänge im Bereich von 680 bis 900 nm in der Farbschicht enthalten sind:

$$R_7 \xrightarrow{Y-N} N=N \xrightarrow{R_3} R_4 R_1$$

$$R_5 \xrightarrow{R_6} R_6$$

$$R_7 \xrightarrow{Y-N} R_2$$

$$R_7 \xrightarrow{Y-N} R_2$$

worin

R₁ und R₂ unabhängig ein Wasserstoffatom, eine Alkylgruppe mit 1 bis 15 Kohlenstoffatomen, eine Arylgruppe mit 6 bis 21 Kohlenstoffatomen, eine Aralkylgruppe mit 7 bis 22 Kohlenstoffatomen oder eine Alkenylgruppe mit 2 bis 16 Kohlenstoffatomen bedeuten:

R₃, R₄, R₅ und R₆ unabhängig ein Wasserstoffatorn, ein Halogenatorn, eine Hydroxylgruppe, eine Carboxylgruppe, eine Sulfonsäuregruppe, eine Sulfonamidgruppe, eine Aminogruppe, eine Alkylgruppe mit 1 bis 15 Kohlenstoffatornen, eine Alkylgruppe mit 1 bis 15 Kohlenstoffatornen, eine Arylgruppe mit 6 bis 21 Kohlenstoffatornen, eine Arylgruppe mit 2 bis 16 Kohlenstoffatornen, eine Alkylcarboxylgruppe mit 2 bis 16 Kohlenstoffatornen, eine Alkylcarbonylaminogruppe mit 2 bis 16 Kohlenstoffatornen, eine Alkylsulfaminogruppe mit 1 bis 15 Kohlenstoffatornen, eine Alkylaminogruppe mit 1 bis 15 Kohlenstoffatornen, eine Alkylsulfaminogruppe mit 1 bis 15 Kohlenstoffatornen oder eine Alkenylgruppe mit 2 bis 16 Kohlenstoffatornen wiedergeben;

R₁ und R₄, R₂ und R₆ und R₁ und R₂ Ringe bilden können;

R₇ ein Wasserstoffatom, ein Halogenatom, eine Hydroxylgruppe, eine Carboxylgruppe, eine Sulfonsäuregruppe, eine Sulfonamidgruppe, eine Aminogruppe, eine Alkylgruppe mit 1 bis 15 Kohlenstoffatomen, eine Alkylgruppe mit 1 bis 15 Kohlenstoffatomen, eine Arylgruppe mit 1 bis 15 Kohlenstoffatomen, eine Arylgruppe mit 2 bis 16 Kohlenstoffatomen, eine Aralkylgruppe mit 7 bis 22 Kohlenstoffatomen, eine Alkylcarboxylgruppe mit 2 bis 16 Kohlenstoffatomen, eine Alkylcarboxylgruppe mit 2 bis 16 Kohlenstoffatomen, eine Alkylcarboxylaminogruppe mit 1 bis 15 Kohlenstoffatomen, eine Alkylcarboxylgruppe mit 1 bis 15 Kohlenstoffatomen, eine Alkylcarboxylgruppe mit 2 bis 16 Kohlenstoffatomen, eine Alkylcarboxylgruppe mit 2 bis 16 Kohlenstoffatomen, eine Cyanogruppe, eine Nitrogruppe, eine Mercaptogruppe, eine Thiocyanogruppe, eine Chlorsulfonsäuregruppe, eine Alkylthiogruppe mit 1 bis 15 Kohlenstoffatomen, eine Alkylcarboxorethingruppe mit 2 bis 16 Kohlenstoffatomen oder eine Alkylcarboxylgruppe mit 1 bis 15 Kohlenstoffatomen bedeutet;

X ein Schwefelatom oder N-R₈ ist, worin R₈ ein Wasserstoffatom, eine Alkylgruppe mit 1 bis 15 Kohlenstoffatomen, eine Arylgruppe mit 6 bis 21 Kohlenstoffatomen, eine Aralkylgruppe mit 7 bis 22 Kohlenstoffatomen oder eine Alkenylgruppe mit 2 bis 16 Kohlenstoffatomen darstellt;

Y für ein Stickstoffatom oder C-R₉ steht, worin R₉ synonym ist mit den für R₇ angegebenen Bedeutungen; mit der Maßgabe, daß, wenn X für ein Schwefelatom steht, Y ein Stickstoffatom bedeutet und, wenn X für N-R₈ steht, Y für C-R₉ steht;

$$(A_{a})_{ma} = (Y_{a})_{1a} = (Y_{$$

worin Y_1 , Y_2 , Y_3 , Y_4 , Y_5 , Y_6 , Y_7 und Y_8 unabhängig ein Wasserstoffatom, eine Kohlenwasserstoffgruppe mit 1 bis 20 Kohlenstoffatomen, eine Alkoxygruppe mit 1 bis 20 Kohlenstoffatomen oder eine Alkylthiogruppe mit 1 bis 20 Kohlenstoffatomen wiedergeben; in den jeweiligen Kombination von Y_1 und Y_2 , Y_3 und Y_4 , Y_5 und Y_6 und Y_7 und Y_8 diese unter Bildung von Ringen kombiniert sein können, wenn sie zueinander benachbart vorliegen;

 A_1 , A_2 , A_3 und A_4 unabhängig ein Halogenatom oder eine Nitrogruppe darstellen; I_1 , I_2 , I_3 und I_4 eine ganze Zahl von 0 bis 3 bedeuten;

m1, m2, m3 und m4 für eine ganze Zahl von 0 bis 3 stehen; und

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M für zwei Wasserstoffatorne, ein zweiwertiges Metallatorn, ein dreiwertiges oder vierwertiges, substituiertes Metallatorn oder ein Oxymetail steht.

 Optisches Aufzeichnungsmedium gemäß Anspruch 1, worin die in der Farbschicht enthaltene Verbindung mit einem Absorptionsmaximum bei einer Wellenlänge im Bereich von 450 bis 630 nm ein Metallkomplex einer Azoverbindung der folgenden Formet (3)

ist, worin R_1 , R_2 , R_4 , R_5 und R_6 synonym sind mit den für Formel (1) angegebenen Bedeutungen; R_{10} eine Hydroxylgruppe oder eine Carboxylgruppe bedeutet; und R_{11} für ein Wasserstoffatom oder ein Halogenatom steht.

- 3. Optisches Aufzeichnungsmedium gemäß Anspruch 1,worin die Farbschicht eine einschichtige Struktur umfaßt, die eine Mischung der Azoverbindung der Formel (1) oder des Metallkomplexes hiervon mit einem Absorptionsmaximum bei einer Wellenlänge im Bereich von 450 bis 630 nm und der Phthalocyaninverbindung der Formel (2) mit einem Absorptionsmaximum bei einer Wellenlänge im Bereich von 680 bis 900 nm enthält.
- 4. Optisches Aufzeichnungsmedium gemäß Anspruch 1,worin die Farbschicht eine zweischichtige Struktur einer optischen Interferenzschicht, enthaltend die Azoverbindung der Formel (1) oder den Metallkomplex hiervon mit einem Absorptionsmaximum bei einer Wellenlänge im Bereich von 450 bis 630 nm,und einer Aufzeichnungsschicht, enthaltend die Phthalocyaninverbindung der Formel (2) mit einem Absorptionsmaximum bei einer Wellenlänge im Bereich von 680 bis 900 nm, umfaßt.
- 5. Optisches Aufzeichnungsmedium gemäß Anspruch 4,worin eine Gleichung von 70≤ni x di≤300 sich auf Licht mit

einer Weltenlänge, die für die Aufzeichnung und Reproduktion verwendet wird, bezieht, worin ni einen realen Teil eines komplexen Brechungsindex bedeutet und di eine Schichtdicke in der optischen Interferenzschicht wiedergibt.

- 6. Optisches Aufzeichnungsmedium gemäß einem der Ansprüche 1 bis 5, das befähigt ist zur Aufzeichnung und/ oder Reproduktion mit einem Laserstrahl mit einer Wellenlänge λ₁, ausgewählt unter Wellenlängen im Bereich von 770 bis 830 nm, und das befähigt ist zur Aufzeichnung und/oder Reproduktion ebenfalls mit einem Laserstrahl mit einer Wellenlänge λ₂, ausgewählt unter Wellenlängen im Bereich von 620 bis 690 nm.
- Optisches Aufzeichnungsmedium gemäß Anspruch 6, das befähigt ist zur Reproduktion mit einem Laserstrahl mit der Wellenlänge \(\lambda_2\).
 - 8. Optisches Aufzeichnungsmedium gemäß Anspruch 6 oder 7, worin der Laserstrahl mit der Wellenlänge λ₁ ein Reflexionsvermögen von 65% oder mehr aufweist und der Laserstrahl mit der Wellenlänge λ₂ ein Reflexionsvermögen von 15% oder mehr aufweist, die durch das Substrat gemessen werden.
 - 9. Optisches Aufzeichnungsmedium gemäß Anspruch 8, worin der Laserstrahl mit der Wellenlänge λ_2 ein Reflexionsvermögen von 20% oder mehr aufweist, das durch das Substrat gemessen wird.
 - 10. Metallkomplex einer Azoverbindung der Formel (4)

$$\begin{bmatrix} R_{11} & N - N \\ S & N = N - N \end{bmatrix} = N \begin{bmatrix} R_{1} \\ R_{2} \end{bmatrix} M e^{2t}$$
(4)

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 $\rm R_1$ und $\rm R_2$ unabhängig ein Wasserstoffatom, eine Alkylgruppe mit 1 bis 15 Kohlenstoffatomen, eine Arylgruppe mit 6 bis 21 Kohlenstoffatomen, eine Aralkylgruppe mit 7 bis 22 Kohlenstoffatomen oder eine Alkenylgruppe mit 2 bis 16 Kohlenstoffatomen bedeuten:

R₁₁ ein Wasserstoffatom oder ein Halogenatom wiedergibt; Me für Nickel, Kobalt, Kupfer, Palladium, Eisen und Zink steht.

Revendications

1. Support d'enregistrement optique comprenant au moins une couche de colorant, une couche réflectrice et une couche protectnce appliquées sur un substrat, dans lequel ladite couche de colorant contient un composé azoïque représenté par la formule (1) ci-dessous ou un de ses complexes métalliques, ayant un maximum d'absorption à une longueur d'onde comprise entre 450 et 630 nm, et un composé de phtalocyanine représenté par la formule (2) ci-dessous ayant un maximum d'absorption à une longueur d'onde comprise entre 680 et 900 nm:

$$R_7 \xrightarrow{Y-N} N=N \xrightarrow{R_3} R_4 \xrightarrow{R_1} R_2 \tag{1}$$

où R₁ et R₂ représentent indépendamment un atome d'hydrogène, un groupe alkyle de 1 à 15 atomes de carbone, un groupe aryle de 6 à 21 atomes de carbone, un groupe aralkyle de 7 à 22 atomes de carbone ou un groupe alcényle de 2 à 16 atomes de carbone ;

 R_3 , R_4 , R_5 et R_6 représentent indépendamment un atome d'hydrogène, un atome d'halogène, un groupe hydroxyle, un groupe acide sulfonique, un groupe sulfonamide, un groupe amino, un groupe alkyle de 1 à 15 atomes de carbone, un groupe alcoxyle de 1 à 15 atomes de carbone, un groupe aryle de 6 à 21 atomes de carbone, un groupe acyle de 1 à 15 atomes de carbone, un groupe alkylcarboxyle de à 16 atomes de carbone, un groupe aralkyle de 7 à 22 atomes de carbone, un groupe alkylcarbonylamino de 2 à 16 atomes de carbone, un groupe alkylsulfoamino de 1 à 15 atomes de carbone, un groupe alkylsulfoamino de 1 à 15 atomes de carbone ou un groupe alkylsulfoamino de 1 à 15 atomes de carbone ou un groupe alkylsulfoamino de 1 à 15 atomes de carbone ou un groupe alcényle de à 16 atomes de carbone;

R₁ et R₄, R₂ et R₆, et R₁ et R₂ peuvent former des cycles ;

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R₇ représente un atome d'hydrogène, un atome d'halogène, un groupe hydroxyle, un groupe carboxyle, un groupe acide sulfonique, un groupe sulfonamide, un groupe amino, un groupe alkyle de 1 à 15 atomes de carbone, un groupe alcoxyle de 1 à 15 atomes de carbone, un groupe acyle de 6 à 21 atomes de carbone, un groupe acyle de 1 à 15 atomes de carbone, un groupe alkylcarboxyle de à 16 atomes de carbone, un groupe aralkyle de 7 à 22 atomes de carbone, un groupe alkylcarbonylamino de 2 à 16 atomes de carbone, un groupe alkylsulfoamino de 1 à 15 atomes de carbone, un groupe alkylsulfonyle de 1 à 15 atomes de carbone, un groupe alcényle de 2 à 16 atomes de carbone, un groupe cyano, un groupe nitro, un groupe mercapto, un groupe thiocyano, un groupe acide chlorosulfonique, un groupe alkylthio de 1 à 15 atomes de carbone, un groupe alkylazométhine de 2 à 16 atomes de carbone ou un groupe alkylaminosulfonyle de 1 à 15 atomes de carbone ou un groupe alkylaminosulfonyle de 1 à 15 atomes de carbone carbone.

X représente un atome de soufre ou N-R₈, où R₈ représente un atome d'hydrogène, un groupe alkyle de 1 à 15 atomes de carbone, un groupe aryle de 6 à 21 atomes de carbone, un groupe aralkyle de 7 à 22 atomes de carbone ou un groupe alcényle de 2 à 16 atomes de carbone :

Y représente un atome d'azote ou C-R₉, où R₉ a la définition décrite pour R₇;

à condition que Y soit un atome d'azote lorsque X est un atome de soufre, et que Y soit C-Rg lorsque X est N-Rg.

$$(A_{4})_{m4} = (Y_{3})_{h4} + (Y_{3})_{m1} + (Y_{3})_{h2} + (Y_{4})_{m2} + (Y_{5})_{m3} + (Y_{$$

où Y_1 , Y_2 , Y_3 , Y_4 , Y_5 , Y_6 , Y_7 et Y_8 représentent indépendamment un atome d'hydrogène, un groupe hydrocarboné de 1 à 20 atomes de carbone, un groupe alkylthio de 1 à 20 atomes de carbone ; dans les combinaisons respectives de Y_1 et Y_2 , Y_3 et Y_4 , Y_5 et Y_6 et Y_7 et Y_8 , ils peuvent être combinés pour former des cycles lorsqu'ils sont adjacents entre eux ;

 A_1 , A_2 , A_3 et A_4 représentent indépendamment un atome d'halogène ou un groupe nitro; I_1 , I_2 , I_3 et I_4 représentent un entier de 0 à 3;

m₁, m₂, m₃ et m₄ représentent un entier de 0 à 3 ; et

M représente deux atomes d'hydrogène, un atome de métal divalent, un atome de métal trivalent ou tétravalent substitué, ou un oxy-métal.

2. Support d'enregistrement optique selon la revendication 1, dans lequel le composé contenu dans la couche de colorant et ayant un maximum d'absorption à une longueur d'onde comprise entre 450 et 630 nm est un complexe métallique d'un composé azolque représenté par la formule (3) ci-dessous :

$$R_{11} \xrightarrow{N-N} N = N \xrightarrow{R_{10}} R_4$$

$$R_{11} \xrightarrow{N-N} N = N \xrightarrow{R_{10}} R_5$$

$$R_{10} \xrightarrow{R_4} R_5$$

$$R_{10} \xrightarrow{R_4} R_5$$

$$R_{10} \xrightarrow{R_4} R_5$$

$$R_{10} \xrightarrow{R_4} R_5$$

dans laquelle R_1 , R_2 , R_4 , R_5 et R_6 ont la définition décrite pour la formule (1); R_{10} représente un groupe hydroxyle ou un groupe carboxyle; et R_{11} représente un atome d'hydrogène ou un atome d'halogène.

- 3. Support d'enregistrement optique selon la revendication 1, dans lequel la couche de colorant comprend une structure monocouche contenant un mélange du composé azolque représenté par la formule (1) ou de son complexe métallique, ayant un maximum d'absorption à une longueur d'onde comprise entre 450 et 630 nm, et du composé de phtalocyanine représenté par la formule (2), ayant un maximum d'absorption à une longueur d'onde comprise entre 680 et 900 nm.
- 4. Support d'enregistrement optique selon la revendication 1, dans lequel la couche de colorant comprend une structure à deux couches constituée d'une couche d'interférence optique contenant le composé azoïque représenté par la formule (1) ou son complexe métallique, ayant un maximum d'absorption à un longueur d'onde comprise entre 450 et 630 nm, et une couche d'enregistrement contenant le composé de phtalocyanine représenté par la formule (2) ayant un maximum d'absorption à une longueur d'onde comprise entre 680 et 900 nm.
 - 5. Support d'enregistrement optique selon la revendication 4, dans lequel l'équation 70 ≤ ni x di ≤ 300 s'applique à la lumière ayant la longueur d'onde utilisée pour l'enregistrement et la reproduction, où ni représente la partie réelle d'un indice de réfraction complexe et di représente l'épaisseur de couche de la couche d'interférence optique.
- 50 6. Support d'enregistrement optique selon l'une quelconque des revendications 1 à 5, pouvant enregistrer et/ou reproduire avec un rayon laser ayant une longueur d'onde λ₁ choisie parmi les longueurs d'onde comprises entre 770 et 830 nm et pouvant aussi enregistrer et/ou reproduire avec un rayon laser ayant une longueur d'onde λ₂ choisie parmi les longueurs d'onde comprises entre 620 et 690 nm.
- Support d'enregistrement optique selon la revendication 6, pouvant reproduire avec un rayon laser ayant la longueur d'onde λ₂.
 - 8. Support d'enregistrement optique selon la revendication 6 ou la revendication 7, dans lequel le rayon laser ayant la longueur d'onde λ₁ a un facteur de réflexion d'au moins 65 %, et le rayon laser ayant la longueur d'onde λ₂ a un facteur de réflexion d'au moins 15 %, ces facteurs de réflexion étant mesurés à travers le substrat.
 - 9. Support d'enregistrement optique selon la revendication 8, dans lequel le rayon laser ayant la longueur d'onde λ_2 a un facteur de réflexion, mesuré à travers le substrat, d'au moins 20 %.
- 45 10. Complexe métallique d'un composé azoïque, représenté par la formule (4)

dans laquelle

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 $\rm B_1$ et $\rm B_2$ représentent indépendamment un atome d'hydrogène, un groupe alkyle de 1 à 15 atomes de carbone, un groupe aryle de 6 à 21 atomes de carbone, un groupe aryle de 6 à 21 atomes de carbone, un groupe aryle de 7 à 22 atomes de carbone ou un groupe alcényle de 2 à 16 atomes de carbone;

R₁₁ représente un atome d'hydrogène ou un atome d'halogène ; Me représente le nickel, le cobalt, le cuivre, le palladium, le fer et le zinc.



Fig.1

